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(21) International Application Number: PCT/US99/11767 (22) International Filing Date: 28 May 1999 (28.05.99) (30) Priority Data: 09/087,479 29 May 1998 (29.05.98) US (71) Applicant: WAYNE HUGHES INSTITUTE [US/US]; 2665 Long Lake Road, St. Paul, MN 55113 (US). (72) Inventors: UCKUN, Fatih, M.; 12590 Ethan Avenue North, White Bear Lake, MN 55110 (US). NARLA, Rama, K.; 1270 Larpenier Avenue, St. Paul, MN 55113 (US). LIU, Xing-Ping; 1010 6th Street S.E., Minneapolis, MN 55414 (US). (74) Agent: DAIGNAULT, Ronald, A.; Merchant & Gould P.C., 3100 Norwest Center, 90 South Seventh Street, Minneapolis, MN 55402-4131 (US).		(81) Designated States: AL, AM, AT, AT (Utility model), AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, CZ (Utility model), DE, DE (Utility model), DK, DK (Utility model), EE, EE (Utility model), ES, FI, FI (Utility model), GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK (Utility model), SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
(54) Title: QUINAZOLINES FOR TREATING BRAIN TUMOR (57) Abstract Novel substituted quinazoline compounds and conjugates useful to inhibit the growth of brain tumor cells and to inhibit adhesion and migration of brain tumor cells. The compounds of the invention include 4-(3'-bromo-4'-hydroxy phenyl)-amino-6,7-dimethoxyquinazoline and this compound covalently bound to EGF.		

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QUINAZOLINES FOR TREATING BRAIN TUMOR

Field of the Invention

This invention relates to novel quinazoline derivatives effective to induce apoptosis of brain tumor cells. In particular, the invention includes novel
5 hydroxy quinazoline derivatives having potent cytotoxicity against human brain tumor cells, including glioblastoma. The novel compounds of the invention further inhibit adhesion of brain tumor cells to extracellular matrix proteins and inhibit migration of brain tumor cells through the extracellular matrix, activities required for tumor metastases.

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Background of the Invention

As the most malignant primary central nervous systems tumors, high grade anaplastic astrocytoma and glioblastoma multiforme respond poorly to contemporary multimodality treatment programs employing surgical resection, radiation therapy and chemotherapy with a median survival of less than one year
15 after initial diagnosis (Pardos, et al., 1997, *Cancer Medicine*, 1:1471-1514; Brandes, et al., 1996, *Cancer Invest.* 14:551-559; Finlay, J.L., 1992, *Pediatric Neuro-Oncology*, 278-297; Pardos, et al., 1998, *Sem. Surgical Oncol.*, 14:88-95). Consequently, the development of effective new agents and novel treatment modalities against these very poor prognosis brain tumors remains a major focal
20 point in translational oncology research.

Glioblastoma multiforme is also a highly invasive primary brain tumor with a disappointingly high local recurrence rate and mortality. New agents capable of inhibiting the infiltration of normal brain parenchyma by glioblastoma cells are urgently needed.

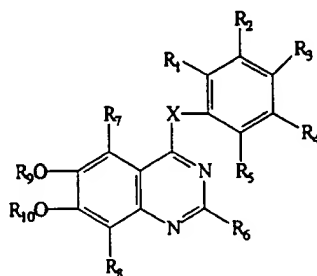
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Summary of the Invention

In a systematic effort to identify a cytotoxic agent with potent anti-tumor activity against glioblastoma cells, several hydroxy-substituted quinazoline-derivatives were synthesized and examined for their *in vitro* and *in vivo* effects on human glioblastoma cells. Novel hydroxy- and halo-hydroxy-quinazoline
30 derivatives were found to exhibit potent cytotoxic activity against human glioblastoma cells at micromolar concentrations. Targeting of these compounds to

the surface of brain tumor cells, for example, by conjugating hydroxy- and the halo-hydroxy compounds to a targeting moiety such as epidermal growth factor (EGF), further enhanced the cytotoxic activity (at nanomolar concentrations). The conjugate demonstrated more rapid and more potent anti-brain tumor activity, including apoptotic death of glioblastoma cells *in vitro*, significantly improved tumor-free survival in an *in vivo* SCID mouse glioblastoma xenograft model, inhibition of tumor cell adhesion to ECM proteins, and inhibition of tumor cell migration and invasion activity.

Accordingly, the present invention includes novel compounds and compositions having potent cytotoxic activity against brain tumor cells. Compositions of the invention contain an effective cytotoxic or inhibitory amount of a hydroxy-substituted quinoxaline compound, more particularly of a hydroxy- or halo-hydroxy-substituted quinazoline derivative. The compounds of the invention include those having the following formula:



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where X is HN, $R_{11}N$, S, O, CH_2 , or $R_{11}CH$, and one or more of R_1 , R_2 , R_3 , R_4 , R_5 is OH, SH, or NH_2 . Preferred embodiments include those where X is HN; R_3 is OH; R_2 and/or R_4 is a halogen, preferably Br. In another preferred embodiment, one or more of R_1 - R_5 form a second ring fused to the phenyl ring, for example, forming a naphthyl ring and having at least one hydroxy substitution.

Preferred cytotoxic compounds of the invention include 4-(3'-Bromo-4'-hydroxylphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P154], 4-(4'-Hydroxylphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P131], and 4-(2'-Hydroxy-naphthyl-3')-amino-6,7-dimethoxyquinazoline [WHI-P292].

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The compounds of the invention can be formulated for delivery to a subject as a pharmaceutical composition, and can preferably be modified for

selective killing of brain tumor cells by conjugation to a cell specific targeting moiety, such as an anti-cell surface antigen-antibody, or a moiety known to bind a cell surface receptor, such as EGF. The compounds of the invention are preferably covalently bonded to the targeting moiety. One exemplary targeting moiety is EGF, which, when conjugated to the compound of the invention, rapidly and specifically directs the compound to brain tumor cells expressing the EGF receptor, resulting in specific, rapid, and enhanced cytotoxicity.

The compounds of the invention are administered to a subject to inhibit the growth of brain tumor cells, to induce apoptosis of brain tumor cells, thereby reducing tumor mass. Compounds of the invention are also administered to inhibit the adhesion and migration of brain tumor cells, for example, inhibiting the infiltration of normal brain parenchyma by glioblastoma cells.

Brief Description of the Drawings

Figures 1A and 1B are cell survival graphs demonstrating the cytotoxic activity of WHI-P154 and other compounds of the invention against U373 glioblastoma cells.

Figures 2A and 2B are photographs of laser analyzed immunostained cells demonstrating cell surface expression of EGF receptor on U373 and U87 human glioblastoma cells.

Figures 3A, 3A', 3B, 3B', 3C, 3C', 3D, and 3D' are photographs of immunostained cells showing EGF-R mediated uptake of EGF-P154 by U373 human glioblastoma cells.

Figure 4A and 4B are photographs showing uptake of WHI-P154 and EGF-P154 by human glioblastoma cells.

Figure 5A is a graph demonstrating the cytotoxic activity of EGF-P154 against brain tumor cells. Figure 5B shows the activity of unconjugated WHI-P154 and EGF-P154 on EGF-R negative leukemia cells.

Figures 6A- 6E are photographs showing morphological features of glioma cells treated with EGF and EGF-P154.

Figures 7A and 7B are graphs showing tumor volume and tumor-free survival times in a SCID mice human glioblastoma model treated with WHI-P154 and EGF-P154.

Figure 8 is a bar graph showing adhesion of glioblastoma and
5 medulloblastoma cells to plates coated with various ECM proteins.

Figures 9A-9D are bar graphs showing inhibition of cell adhesion to ECM proteins in the presence of the compounds of the invention.

Figure 10 is a graph showing the inhibition of EGF-induced cell adhesion in the presence of the compounds of the invention.

10 Figures 11A and 11B are graphs demonstrating inhibition of brain tumor cell migration through the extracellular matrix MATRIGEL Matrix in the presence of the compounds of the invention.

Figures 12A-C and 12 A'-C' are photographs of immunostained cells demonstrating the inhibition of focal adhesion complex formation in glioblastoma
15 cells in the presence of the compounds of the invention.

Figures 13A-13C are photographs of immunostained cells demonstrating the inhibition of actin stress fiber formation in glioblastoma cells contacted with the compounds of the invention.

Figures 14A-D are photographs showing inhibition of tumor cell
20 outgrowth from glioblastoma spheroids in the presence of the compounds of the invention.

Figure 15 is a graph demonstrating the cytotoxic effect of WHI-P292 on glioblastoma cells.

Detailed Description of the Invention

25 The present invention includes novel hydroxy-substituted quinazoline derivatives having potent activity as cytotoxic agents against brain tumor cells, including glioblastoma cells. In addition, the hydroxy-substituted quinazoline compounds of the invention are potent inhibitors of tumor cell adhesion and migration, activities required for tumor cell metastases.

Definitions

All scientific and technical terms used in this application have meanings commonly used in the art unless otherwise specified. As used in this application, the following words or phrases have the meanings specified.

5 As used herein, "alkyl" includes both branched and straight-chain saturated aliphatic hydrocarbon groups having the specified number of carbon atoms. As a preferred embodiment, chains of 1 to 4 carbon atoms are included, for example methyl, ethyl, propyl, isopropyl, butyl, secondary butyl, t-butyl, and the like.

10 As used herein, "alkene" includes both branched and straight-chain unsaturated aliphatic hydrocarbon groups having the specified number of carbon atoms, preferably chains of 1 to 4 carbon atoms.

As used herein, "acyl" includes $-C(O)R$, where R is H, alkyl, or aryl containing 1 to 4 carbon atoms.

15 As used herein "halogen" includes fluoro, chloro, bromo, and iodo. A preferred halogen or halo substituent is Br.

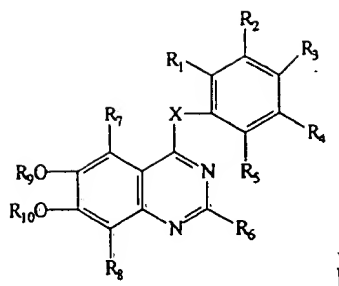
20 As used herein, "pharmaceutically acceptable carrier" includes any material which, when combined with a compound of the invention, allows the compound to retain biological activity, such as the ability to induce apoptosis of brain tumor cells, and is non-reactive with the subject's immune system. Examples include, but are not limited to, any of the standard pharmaceutical carriers such as a phosphate buffered saline solution, water, emulsions such as oil/water emulsions, and various types of wetting agents. Compositions comprising such carriers are formulated by well known conventional methods (see, for example, Remington's Pharmaceutical Sciences, Chapter 43, 14th Ed., Mack Publishing Co., Easton, PA).

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Compounds of the Invention

The novel substituted quinazolines of the invention have the general structure represented by the following formula I:

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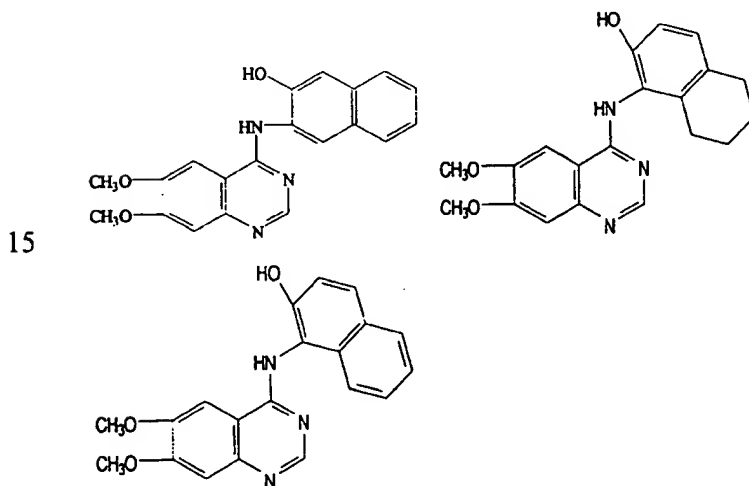


where X is selected from the group consisting of HN, $R_{11}N$, S, O, CH_2 , and $R_{11}-CH$. R_{11} is H, alkyl, having 1 to 4 carbon atoms, or acyl. Preferably, X is NH; and preferably R_{11} is H.

- 5 $R_1, R_2, R_3, R_4, R_5, R_6, R_7$ and R_8 are each independently selected from the group consisting of H, OH, SH, NH_2 , NO_2 , alkoxy, alkylthio and halogen. R_9 and R_{10} are each independently selected from the group consisting of H, alkyl or acyl, containing up to 4 carbon atoms. Preferably, R_9 and R_{10} are methyl.

- At least one of R_1, R_2, R_3, R_4, R_5 is OH. Alternatively, at least one of
10 $R_1 - R_5$ is a compound such as SH or NH_2 .

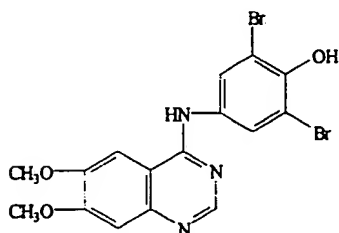
In an alternative embodiment, one or more of R_1-R_5 forms a second ring fused to the phenyl ring. For example, the following compounds include second rings fused to the phenyl ring via one or more of R_1-R_5 :



Exemplary compounds

Some exemplary compounds of the invention are listed below with their characterization data:

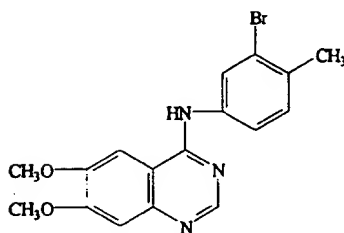
4-(3',5'-Dibromo-4'-hydroxyphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P97]



yield 72.80%; m.p. >300.0 °C. UV(MeOH) λ_{max} : 208.0, 210.0, 245.0, 320.0 nm; IR(KBr) ν_{max} : 3504(br), 3419, 2868, 1627, 1512, 1425, 1250, 1155 cm^{-1} ; ^1H NMR(DMSO- d_6): δ 9.71(s, 1H, -NH), 9.39(s, 1H, -OH), 8.48(s, 1H, 2-H), 8.07(s, 2H, 2', 6'-H), 7.76(s, 1H, 5-H), 7.17(s, 1H, 8-H), 3.94(s, 3H, -OCH₃), 3.91(s, 3H, -OCH₃). GC/MS m/z 456(M^+ +1, 54.40), 455(M^+ , 100.00), 454(M^+ -1, 78.01), 439(M^+ -OH, 7.96), 376(M^+ +1 - Br, 9.76), 375(M^+ -Br, 10.91), 360(5.23). Anal. (C₁₆H₁₃Br₂N₃O₃) C, H, N.

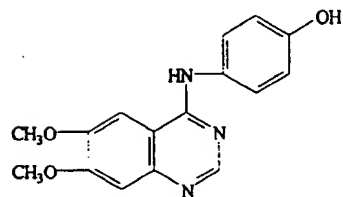
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4-(3'-Bromo-4'-methylphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P111]

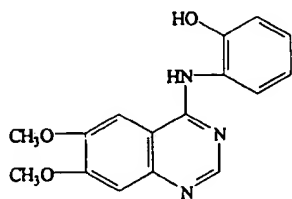


yield 82.22 %; m.p. 225.0-228°C. ^1H NMR(DMSO- d_6): δ 10.23(s, 1H, -NH), 8.62(s, 1H, 2-H), 8.06(d, 1H, $J_{2,6}=2.1$ Hz, 2'-H), 7.89(s, 1H, 5-H), 7.71(dd, 1H, $J_{5,6}=8.7$ Hz, $J_{2,6}=2.1$ Hz, 6'-H), 7.37(d, 1H, $J_{5,6}=8.7$ Hz, 5'-H), 7.21(s, 1H, 8-H), 3.96(s, 3H, -OCH₃), 3.93(s, -OCH₃). UV(MeOH) λ_{max} (ϵ): 204.0, 228.0, 255.0, 320.0 nm. IR(KBr) ν_{max} : 3431, 3248, 2835, 1633, 1517, 1441, 1281, 1155 cm^{-1} . GC/MS m/z 375(M^+ +1, 76.76), 374(M^+ , 100.00), 373(M^+ -1, 76.91), 358(M^+ +1 - OH, 11.15), 357(1.42), 356(6.31). Anal. (C₁₇H₁₆BrN₃O₂HCl) C, H, N.

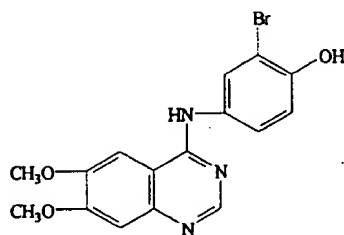
4-(4'-Hydroxyphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P131]



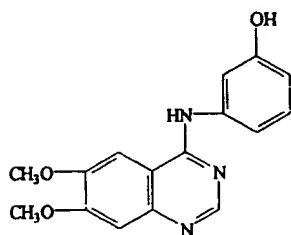
yield 84.29%; m.p. 245.0-248.0 °C. UV(MeOH) λ_{max} : 203.0, 222.0, 251.0, 320.0 nm; IR(KBr) ν_{max} : 3428, 2836, 1635, 1516, 1443, 1234 cm^{-1} ; ^1H NMR(DMSO- d_6): δ 11.21(s, 1H, -NH), 9.70(s, 1H, -OH), 8.74(s, 1H, 2-H), 8.22(s, 1H, 5-H), 7.40(d, 2H, $J=8.9$ Hz, 2', 6'-H), 7.29(s, 1H, 8-H), 6.85(d, 2H, $J=8.9$ Hz, 3', 5'-H), 3.98(s, 3H, -OCH₃), 3.97(s, 3H, -OCH₃). GC/MS m/z 298(M^+ +1, 100.00), 297(M^+ , 26.56), 296(M^+ -1, 12.46). Anal. (C₁₆H₁₅N₃O₃HCl) C, H, N.

4-(2'-Hydroxyphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P132]

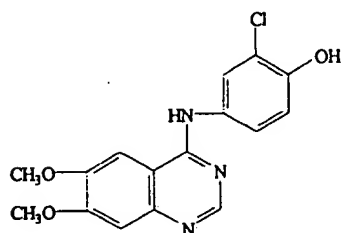
yield 82.49%; m.p. 255.0-258.0 °C. ¹H NMR(DMSO-d₆): δ 9.78(s, 1H, -NH), 9.29(s, 1H, -OH), 8.33(s, 1H, 2-H), 7.85(s, 1H, 5-H), 7.41-6.83(m, 4H, 3', 4', 5', 6'-H), 7.16(s, 1H, 8-H), 3.93(s, 3H, -OCH₃), 3.92(s, 3H, -OCH₃). UV(MeOH)λ_{max}(ε): 203.0, 224.0, 245.0, 335.0 nm. IR(KBr)ν_{max}: 3500 (br), 3425, 2833, 1625, 1512, 1456, 1251, 1068 cm⁻¹. GC/MS m/z 298(M⁺ +1, 8.91), 297(M⁺, 56.64), 281(M⁺ +1 - OH, 23.47), 280(M⁺ - OH, 100.00). Anal. (C₁₆H₁₅N₃O₃HCl) C, H, N.

4-(3'-Bromo-4'-hydroxyphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P154]

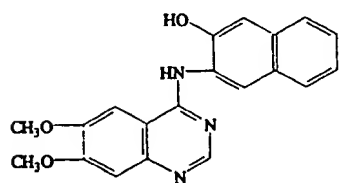
yield 89.90%; m.p. 233.0-233.5 °C. UV(MeOH)λ_{max}: 203.0, 222.0, 250.0, 335.0 nm; IR(KBr) ν_{max}: 3431 br, 2841, 1624, 1498, 1423, 1244 cm⁻¹; ¹H NMR(DMSO-d₆): δ 10.08(s, 1H, -NH), 9.38(s, 1H, -OH), 8.40(s, 1H, 2-H), 7.89(d, 1H, J_{2',5'} = 2.7 Hz, 2'-H), 7.75(s, 1H, 5-H), 7.55(dd, 1H, J_{5',6'} = 9.0 Hz, J_{2',6'} = 2.7 Hz, 6'-H), 7.14(s, 1H, 8-H), 6.97(d, 1H, J_{5',6'} = 9.0 Hz, 5'-H), 3.92(s, 3H, -OCH₃), 3.90(s, 3H, -OCH₃). GC/MS m/z 378(M⁺ +2, 90.68), 377(M⁺ +1, 37.49), 376(M⁺, 100.00), 360(M⁺, 3.63), 298(18.86), 282 (6.65). Anal. (C₁₆H₁₄N₃O₃HCl) C, H, N.

4-(3'-Hydroxyphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P180]

yield 71.55 %; m.p. 256.0- 258.0 °C. ¹H NMR(DMSO-d₆): δ 9.41(s, 1H, -NH), 9.36(s, 1H, -OH), 8.46(s, 1H, 2-H), 7.84(s, 1H, 5-H), 7.84-6.50(m, 4H, 2', 4', 5', 6'-H), 7.20(s, 1H, 8-H), 3.96(s, 3H, -OCH₃), 3.93(s, 3H, -OCH₃). UV(MeOH)λ_{max}(ε): 204.0, 224.0, 252.0, 335.0 nm. IR(KBr)ν_{max}: 3394, 2836, 1626, 1508, 1429, 1251 cm⁻¹. GM/MS m/z: 297(M⁺, 61.89), 296(M⁺, 61.89), 296(M⁺ -1, 100.00), 280(M⁺ - OH, 13.63). Anal. (C₁₆H₁₅N₃O₃.HCl) C, H, N.

4-(3'-Chloro-4'-Hydroxyphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P197]

yield 84.14%; m.p. 245.0°C(dec). ¹H NMR(DMSO-d₆): δ 10.00(s, 1H, -NH), 9.37(s, 1H, -OH), 8.41(s, 1H, 2-H), 7.78(s, 1H, 5-H), 7.49(d, 1H, J_{2,6}=2.7 Hz, 2'-H), 7.55(dd, 1H, J_{5,6}=9.0 Hz, J_{2,6}=2.7 Hz, 6'-H), 7.16(s, 1H, 8-H), 6.97(d, 1H, J_{5,6}=9.0 Hz, 5'-H), 3.93(s, 3H, -OCH₃), 3.91(s, 3H, -OCH₃). UV(MeOH)λ_{max}(ε): 209.0, 224.0, 249.0, 330.0 nm. IR(KBr)ν_{max}: 3448, 2842, 1623, 1506, 1423, 1241 cm⁻¹. GC/MS m/z: 341(M⁺, 100.00), 326(M⁺-CH₃, 98.50), 310(M⁺-OCH₃, 12.5), 295(9.0), 189(13.5), 155(13.8). Anal. (C₁₆H₁₄ClN₃O₃·HCl) C, H, N.

4-(2'-Hydroxy-naphthyl-3')-amino-6,7-dimethoxyquinazoline [WHI-P292].

Yield 87.41%; m.p. 277.0-279.0°C, IR(KBr)ν_{max}: 3479, 3386, 3036, 2901, 1632, 1581, 1504, 1437, 1281 cm⁻¹. ¹H NMR(DMSO-d₆): δ 11.38(s, 1H, -NH), 10.35(s, 1H, -OH), 8.73(s, 1H, 2-H), 8.25(s, 1H, 5-H), 7.93-7.30(m, 6H, 1', 4', 5', 6', 7', 8'-H), 7.37(s, 1H, 8H), 4.00(s, 6H, -OCH₃). GC/MS m/a: 281(41.0), 253(11.0), 207(100.0). Anal. (C₂₀H₁₇N₃O₃·HCl) C, H, N.

Cytotoxic compounds

- 5 As shown in the Examples below, a hydroxyl substituent on the phenyl ring (R₁ - R₅) appears to be necessary for cytotoxic effects of the novel substituted compounds of the invention, while a second substitution in this ring, e.g., with a halogen such as bromine (Br), enhances the cytotoxic effect of the compound. In the method of the invention, the cytotoxic effects of these compounds is achieved
- 10 by contacting brain tumor cells with micromolar amounts of the inhibitory compound. Particularly useful cytotoxic compounds include WHI-P154 (3-Br, 4-OH substituted) and WHI-P131 (4-OH substituted), and WHI-P292 (1-OH naphthalene-substituted). More particularly useful are conjugates of these compounds with a targeting moiety such as EGF, having cytotoxic activity at nanomolar
- 15 concentrations.

As described above, compounds useful in the method of the invention also include those substituted with SH or NH₂ in place of the demonstrated hydroxy substitutions.

5 **Compounds for Inhibiting Adhesion/Migration**

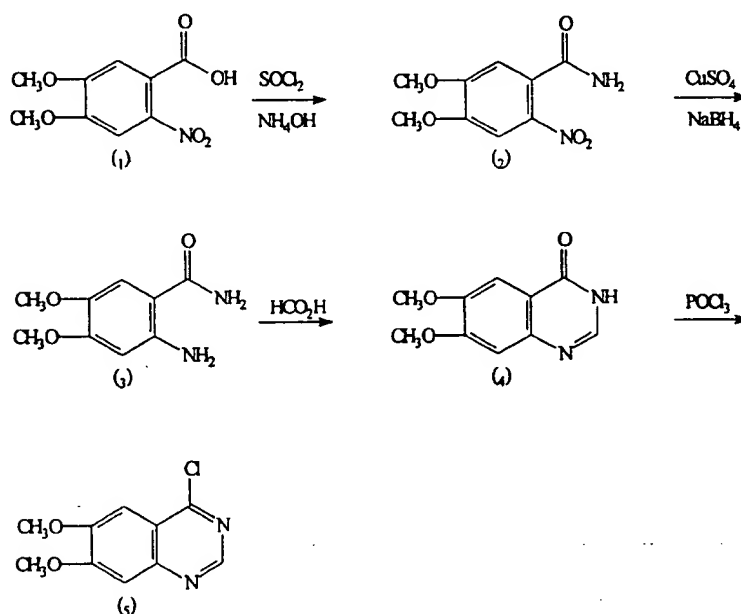
The Examples below further demonstrate the effectiveness of the hydroxy-substituted quinazoline compounds of the invention as inhibitors of brain tumor cell adhesion to extracellular matrix and of tumor cell migration. Each of the tested compounds having a hydroxy substituents on the phenyl ring demonstrated
10 inhibitory activity against glioblastoma cell adhesion/migration. Particularly potent and useful inhibitory compounds include WHI-P154, WHI-P131, and WHI-P292.

Useful compounds of the invention are tested for the ability to prevent adhesion/migration of brain tumor cells by assays described in the Examples below. Such assays include inhibition of cell binding to extracellular matrix proteins
15 in the presence of the inhibitory compound as compared with a non-inhibitory control; inhibition of brain tumor cell invasion into Matrigel Matrix according to the method published by Albini et.al., 1987, *Cancer Res.* 47:3239; and inhibition of focal adhesion plaques and actin polymerization in the presence of the inhibitory compound as compared with a non-inhibitory control. Conjugation of the inhibitory
20 compounds to a targeting moiety, EGF, enhanced the inhibitory activity of the compound WHI-P154.

In the method of the invention, brain tumor cells are contacted with approximately micromolar concentrations of the inhibitory compounds to inhibit tumor cell adhesion and invasion/migration into non-diseased tissue. This is
25 important, for example, during ablation surgery when cells may be dispersed. Adhesion of cells to ECM coupled with the aggressive malignant nature of brain tumor cells can result in new tumor growth at the adhesion site. Inhibition of adhesion and migration by administering the compounds of the invention thereby inhibits new tumor growth.

Synthesis of Novel Hydroxy-Substituted Quinazoline Derivatives

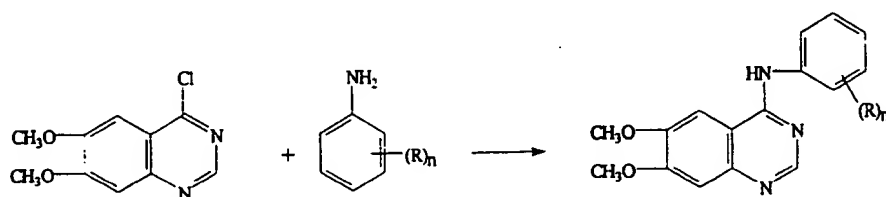
The hydroxy-substituted quinazoline derivatives of the invention can be synthesized from a key starting material, 4-chloro-6,7-dimethoxyquinazoline, prepared using published procedures (Nomoto, et al., 1990, *Chem. Pharm. Bull.*, 38:1591-1595; Thomas, C. L., 1970, *Academic Press, New York, NY*, "I. Synthesis of quinazoline derivatives") as outlined below in Scheme 1 and as described more fully in the Examples below:



10

Scheme 1

The compounds of the invention are then prepared by the condensation of 4-chloro-6,7-dimethoxyquinazoline with the appropriate substituted aniline as outlined below in Scheme 2:

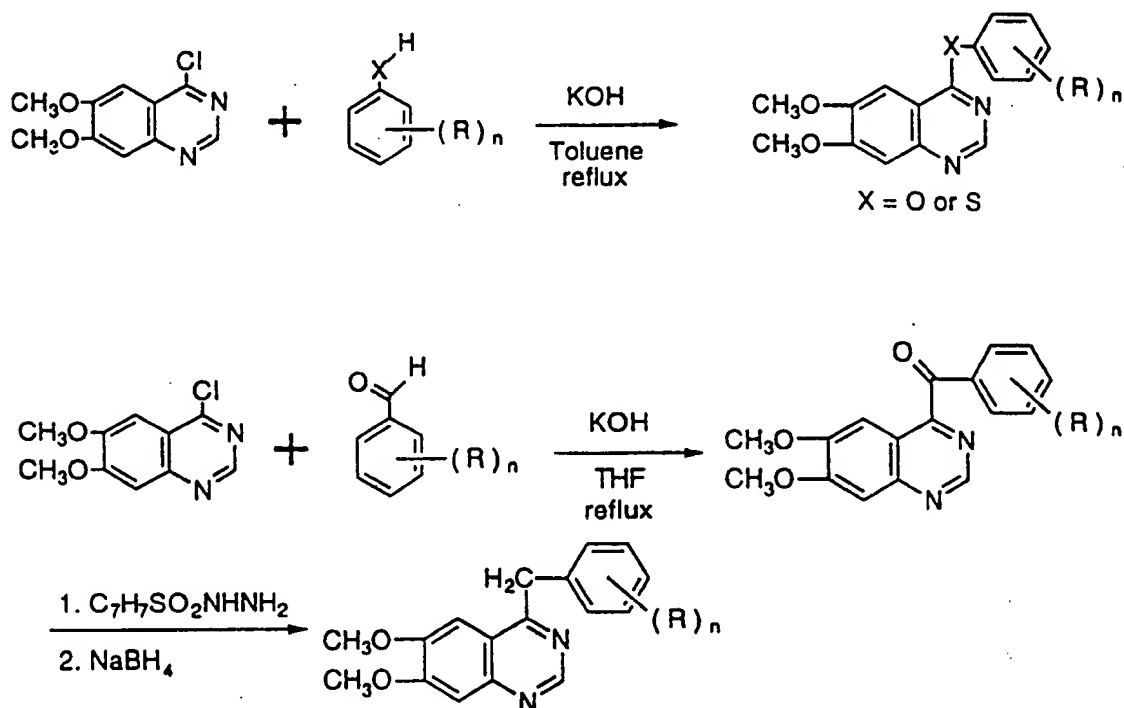


15

R = substituent; n = number

Scheme 2

In a similar manner, compounds of the invention where X is S or O, or where X is alkyl, such as CH₂, are synthesized from a precursor compound, reacting the ring moiety with the desired substitution to produce the desired product. These synthetic methods are known to those in the art, and include, for example, those shown in Scheme 3 below.



20

Scheme 3

Conjugates of the Invention

The term "conjugate" is meant to include a compound formed as a composite between two or more molecules. More specifically, in the present invention, the novel hydroxy-substituted quinazoline derivatives are bonded, for example, covalently bonded, to cell-specific targeting moieties forming a conjugate compound for efficient and specific delivery of the agent to a cell of interest.

Targeting Moiety

The phrase "targeting moiety" is meant to include a molecule which serves to deliver the compounds of the invention to a specific site for the desired activity. Targeting moieties include, for example, molecules that specifically bind

30

molecules on a specific cell surface. Such targeting moieties useful in the invention include anti-cell surface antigen antibodies, growth factors which bind to cell surface receptors such as EGF and its receptor EGF-R. Cytokines, including interleukins and factors such as granulocyte/macrophage stimulating factor (GMCSF) are also specific targeting moieties, known to bind to specific cells expressing high levels of their receptors.

Epidermal Growth Factor (EGF) and Its Receptor (EGF-Rc)

Human Epidermal Growth Factor (hEGF) is commercially available in a highly purified form, for example, from Upstate Biotechnology, Inc. (Lake Placid, NY) (Lot No. 01-107C). This protein ligand is known to bind specifically and with high affinity to receptors located on the surface of EGF-responsive cells. Expression of the EFG-Rc is increased in EGF-responsive cells, including hyperplastic neointima cells.

For use in the conjugates of the present invention, recombinant human EGF (hrEGF) is preferred, although it is anticipated that hEGF and hEGF analogs that specifically bind hEGF-Rc on neointima cells will similarly inhibit migration of vascular cells and formation of hyperplastic neointima cell growth when conjugated.

Human epidermal growth factor (EGF) is a 53 amino acid, single-chain, 6216 daltons polypeptide, which exerts biologic effects by binding to a specific 170 kDa cell membrane epidermal growth factor receptor (EGF-receptor/ErbB-1) (Fix, S.B., 1994, *Breast Cancer Research & Treatment*, 29:41-49; Earp et al., 1995, *Breast Cancer Research & Treatment*, 35:115-132; Wright, et al., 1995, *J. Biol. Chem.*, 270:12085-12093; Broome and Hunter, 1996, *J. Biol. Chem.*, 271:16798-16806). The human EGF-receptor consists of an extracellular domain with a high cysteine content and N-linked glycosylation, a single transmembrane domain, and a cytoplasmic domain with protein tyrosine kinase (PTK) activity.

Binding of EGF to the EGF-receptor/ErbB-1 results in receptor dimerization with itself or other members of the Erb-B (subtype I) transmembrane PTK family (e.g., Erb-B2, Erb-B3), resulting in activation with autophosphorylation of the PTK domain (Muthuswamy, S.K., 1994, *Molecular & Cellular Biology*,

14:735-743); Ottenhoff-Kalff et al., 1992, *Cancer Research*, 52:4773-4778). The EGF-receptor is physically and functionally associated with Src protooncogene family PTK, including p60^{src} (Muthuswamy, S.K., 1994, *Molecular & Cellular Biology*, 14:735-743; Ottenhoff-Kalff et al., 1992, *Cancer Research*, 52:4773-4778; 5 Aikyama et al., 1987, *J. Biol. Chem.*, 262:5592-5595). This association is believed to be an integral part of the signaling events mediated by the EGF-receptor (Ottenhoff-Kalff et al., 1992, *Cancer Research*, 52:4773-4778).

Conjugate formation

10 To form the conjugates of the invention, targeting moieties are covalently bonded to sites on the hydroxy-substituted quinazoline compounds. The targeting moiety, which is often a polypeptide molecule, is bound to compounds of the invention at reactive sites, including NH₂, SH, CHO, COOH, and the like. Specific linking agents are used to link the compounds. Preferred linking agents are 15 chosen according to the reactive site to which the targeting moiety is to be attached.

Methods for selecting an appropriate linking agent and reactive site for attachment of the targeting moiety to the compound of the invention are known, and are described, for example, in Hermanson, et al., *Bioconjugate Techniques*, Academic Press, 1996; Hermanson, et al., *Immobilized Affinity Ligand Techniques*, 20 Academic Press, 1992; and *Pierce Catalog and Handbook*, 1996, pp. T155-T201. One exemplary method for conjugating EGF is described in the Examples below.

Administration Methods

The conjugates of the present invention can be formulated as 25 pharmaceutical compositions and administered to a mammalian host, including a human patient in a variety of forms adapted to the chosen route of administration and suitable for administration of the small molecule or its conjugate. Preferred administration routes include orally, parenterally, as well as intravenous, intramuscular or subcutaneous routes.

30 It is preferred that the conjugate of the present invention be administered parenterally, i.e., intravenously or intraperitoneally, by infusion or injection. In one embodiment of the invention, the compounds may be administered

directly to a tumor by tumor injection; by injecting the compound into the brain, e.g., into the ventricular fluid; or by systemic delivery by intravenous injection. The compounds of the invention, including the conjugates, are of a size and composition expected to have ready access to the brain across the blood-brain barrier.

5 Solutions or suspensions of the conjugates can be prepared in water, isotonic saline (PBS) and optionally mixed with a nontoxic surfactant. Dispersions may also be prepared in glycerol, liquid polyethylene, glycols, DNA, vegetable oils, triacetin and mixtures thereof. Under ordinary conditions of storage and use, these preparations may contain a preservative to prevent the growth of microorganisms.

10 The pharmaceutical dosage form suitable for injection or infusion use can include sterile, aqueous solutions or dispersions or sterile powders comprising an active ingredient which are adapted for the extemporaneous preparation of sterile injectable or infusible solutions or dispersions. In all cases, the ultimate dosage form should be sterile, fluid and stable under the conditions of manufacture and
15 storage. The liquid carrier or vehicle can be a solvent or liquid dispersion medium comprising, for example, water, ethanol, a polyol such as glycerol, propylene glycol, or liquid polyethylene glycols and the like, vegetable oils, nontoxic glyceryl esters, and suitable mixtures thereof. The proper fluidity can be maintained, for example, by the formation of liposomes, by the maintenance of the required particle size, in
20 the case of dispersion, or by the use of nontoxic surfactants. The prevention of the action of microorganisms can be accomplished by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, sorbic acid, thimerosal, and the like. In many cases, it will be desirable to include isotonic agents, for example, sugars, buffers, or sodium chloride. Prolonged absorption of
25 the injectable compositions can be brought about by the inclusion in the composition of agents delaying absorption—for example, aluminum monostearate hydrogels and gelatin.

 Sterile injectable solutions are prepared by incorporating the conjugates in the required amount in the appropriate solvent with various other
30 ingredients as enumerated above and, as required, followed by filter sterilization. In the case of sterile powders for the preparation of sterile injectable solutions, the preferred methods of preparation are vacuum drying and freeze-drying techniques,

which yield a powder of the active ingredient plus any additional desired ingredient present in the previously sterile-filtered solutions.

Tumor Treatment

5 For purposes of this invention, a method of tumor treatment includes contacting brain tumor cells with a compound of the invention in order to achieve an inhibition of tumor cell growth, a killing of tumor cells, and/or increased patient survival time. Treatment of tumors, by the method of the invention, also includes the prevention of the adhesion and migration of tumor cells, thereby inhibiting
10 metastases.

 The cytotoxic and adhesion/migration-inhibiting compounds of the invention are suitable for use in mammals. As used herein, "mammals" means any class of higher vertebrates that nourish their young with milk secreted by mammary glands, including, for example, humans, rabbits, and monkeys.

15

Apoptosis

 Apoptosis, or programmed cellular death, is an active process requiring new protein synthesis. Typically, the process requires ATP, involves new RNA and protein synthesis, and culminates in the activation of endogenous
20 endonucleases that degrade the DNA of the cell, thereby destroying the genetic template required for cellular homostasis. Apoptosis is observed in controlled deletion of cells during metamorphosis, differentiation, and general cell turnover and appears normally to be regulated by receptor-coupled events. For these reasons, apoptosis has been called "programmed cell death" or "cell suicide." While every
25 cell likely has the genetic program to commit suicide, it is usually suppressed. Under normal circumstances, only those cells no longer required by the organism activate this self-destruction program.

 Apoptotic cell death is characterized by plasma membrane blebbing, cell volume loss, nuclear condensation, and endonucleolytic degradation of DNA at
30 nucleosome intervals. Loss of plasma membrane integrity is a relatively late event in apoptosis, unlike the form of cell death termed necrosis, which can be caused by

hypoxia and exposure to certain toxins and which is typically characterized early-on by increased membrane permeability and cell rupture.

Adhesion/Migration

5 Adhesion is meant to include that activity of a cell, such as a tumor cell, by which it adheres to extracellular matrix proteins, including laminin, fibronectin, and collagen. Adhesion assays are known, and include, for purposes of this invention, the adherence of tumor cells to plates coated with extracellular matrix proteins.

10 Migration is meant to include that activity of tumor cells by which they migrate through extracellular matrix and invade tissues. Assays for migration include the ability of cells to migrate through a matrix formed of extracellular matrix, such as MATRIGEL matrix, as well as evaluation of the cell's cytoskeletal organization including actin cytoskeletal rearrangement and changes in focal
15 adhesions as described in the following Examples.

Useful Dose

When used *in vivo* to selectively kill brain tumor cells or to inhibit brain tumor cell adhesion/migration, the administered dose is that effective to have
20 the desired effect, e.g., sufficient to reduce or eliminate brain tumors, or sufficient to inhibit adherence/migration of tumor cells. Appropriate amounts can be determined by those skilled in the art, extrapolating using known methods and relationships, from the *in vitro* and *in vivo* data provided in the Examples. Based on the SCID mouse pharmacology data contained in this application, effective exposure levels are
25 expected to be achieved.

In general, the dose of the novel substituted quinozalines effective to achieve brain tumor cell apoptosis, reduction in tumors, and increased survival time, is that which administers micromolar amounts of the compound to the cells, preferably 100 micromolar or greater. The required dose is lessened by conjugation
30 of the compound to a targeting moiety, for example, to preferably 100 nanomolar or greater concentrations.

For cell adhesion and migration inhibitory activities, the compound is administered generally at lower dosages, in the range of 100 micromolar or less.

The effective dose to be administered will vary with conditions specific to each patient. In general, factors such as the disease burden, tumor
5 location (exposed or remote), host age, metabolism, sickness, prior exposure to drugs, and the like contribute to the expected effectiveness of a drug. One skilled in the art will use standard procedures and patient analysis to calculate the appropriate dose, extrapolating from the data provided in the Examples.

10 In general, a dose which delivers about 1-100 mg/kg body weight is expected to be effective, although more or less may be useful.

In addition, the compositions of the invention may be administered in combination with other anti-tumor therapies. In such combination therapy, the administered dose of the hydroxy-substituted quinazoline derivatives would be less than for single drug therapy.

15

EXAMPLES

The invention may be further clarified by reference to the following Examples, which serve to exemplify some of the preferred embodiments, and not to limit the invention in any way.

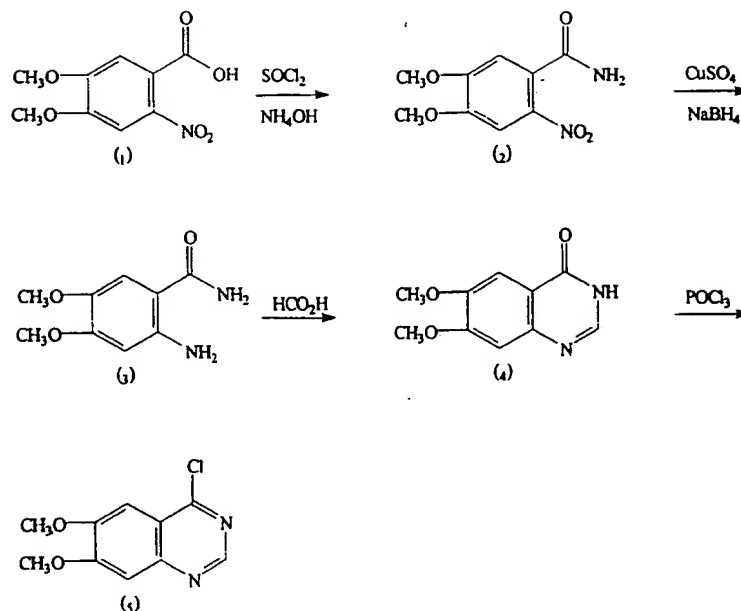
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Example 1

Synthesis of Quinazoline Derivatives

All chemicals were purchased from the Aldrich Chemical Company,
25 Milwaukee, Wisconsin, and were used directly for synthesis. Anhydrous solvents such as acetonitrile, methanol, ethanol, ethyl acetate, tetrahydrofuran, chloroform, and methylene chloride were obtained from Aldrich as sure seal bottles under nitrogen and were transferred to reaction vessels by cannulation. All reactions were carried out under a nitrogen atmosphere.

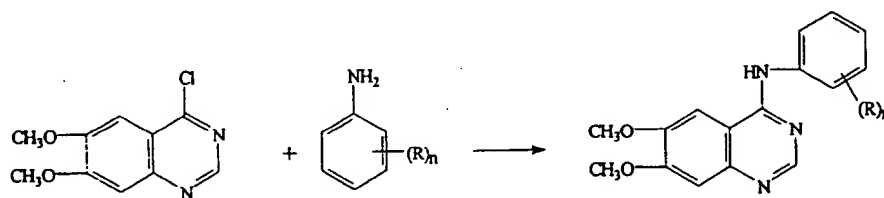
30 The key starting material, 4-chloro-6,7-dimethoxyquinazoline, was prepared using published procedures (Nomoto, et al., 1990, *Chem. Pharm. Bull.*, 38:1591-1595; Thomas, C. L., 1970, *Academic Press, New York, NY*, "I. Synthesis of quinazoline derivatives") as outlined below in Scheme 1:



Scheme 1

Specifically, 4,5-dimethoxy-2-nitrobenzoic acid (compound 1) was treated with thionyl chloride to form acid chloride, followed by reacting with ammonia to yield 4,5-dimethoxy-2-nitrobenzamide (compound 2). Compound 2 was reduced with sodium borohydride in the presence of catalytic amounts of copper sulphate to give 4,5-dimethoxy-2-aminobenzamide (compound 3), which was directly refluxed with formic acid to yield 6,7-dimethoxyquinazoline-4(3H)-one (compound 4). Compound 4 was refluxed with phosphorus oxytrichloride to give 4-chloro-6,7-dimethoxyquinazoline (compound 5) in good yield.

Substituted quinazoline derivatives were prepared by the condensation of 4-chloro-6,7-dimethoxyquinazoline with substituted anilines as outlined below in Scheme 2:

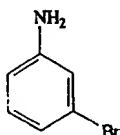
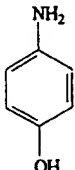
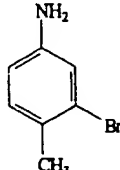
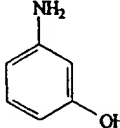
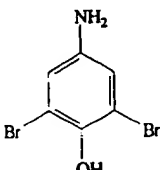
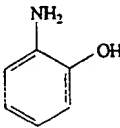
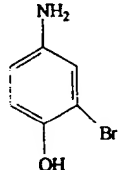
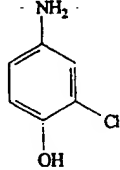
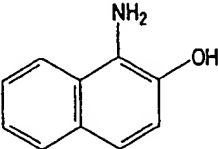
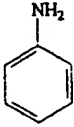


R = substituent; n = number

Scheme 2

Specifically, a mixture of 4-chloro-6,7-dimethoxyquinazoline (448 mg, 2 mmols) and the substituted aniline (2.5 mmols) in EtOH (20 ml) was heated to reflux. After refluxing for 4-24 hours, an excess amount of Et₃N was added, and the solvent was concentrated to give the crude product which was recrystallized from DMF.

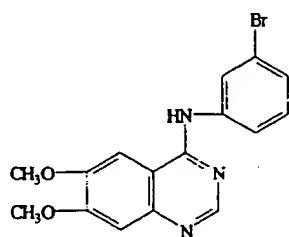
As discussed above, the novel hydroxy-substituted quinazoline derivatives of the invention were created by reacting substituted anilines with the key starting material, 4-chloro-6,7-dimethoxyquinazoline. Each of the anilines to synthesize the compounds is shown in the table below.

forms WHI-P79 	forms WHI-P131 	forms WHI-P111 	forms WHI-P180 
forms WHI-P97 	forms WHI-P132 	forms WHI-P154 	forms WHI-P197 
forms WHI-P292 			forms WHI-P258 

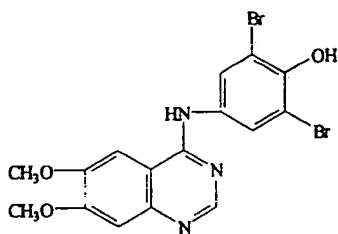
Example 2
Characterization of Substituted Quinazoline Derivatives

5 The substituted quinazoline derivatives were synthesized as described in Example 1 and characterized. Each structure is shown below, along with its identifying analytical test results. Proton and carbon Nuclear Magnetic Resonance (^1H and ^{13}C NMR) spectra were recorded on a Mercury 2000 Varian spectrometer operating at 300 MHz and 75 MHz, respectively, using an automatic broad band
10 probe. Unless otherwise noted, all NMR spectra were recorded in CDCl_3 at room temperature. ^1H chemical shifts are quoted in parts per million (δ in ppm) downfield from tetramethyl silane (TMS), which was used as an internal standard at 0 ppm and s, d, t, q, m designate singlet, doublet, triplet, quartet and multiplet, respectively. Melting points were determined using a Fisher-Johns melting apparatus and are
15 uncorrected. UV spectra were recorded using a Beckmann Model # DU 7400 UV/V is spectrometer with a cell path length of 1 cm. Methanol was used as the solvent for the UV spectra. Fourier Transform Infrared spectra were recorded using an FT-Nicolet model Protege #460 instrument. The infrared spectra of the liquid samples were run as neat liquids using KBr discs. The KBr pellet method was used for all
20 solid samples. The GC/mass spectrum analysis was conducted using a Hewlett-Packard GC/mass spectrometer model # 6890 equipped with a mass ion detector and Chem Station software. The temperature of the oven was steadily increased from 70°C to 250°C and the carrier gas was helium.

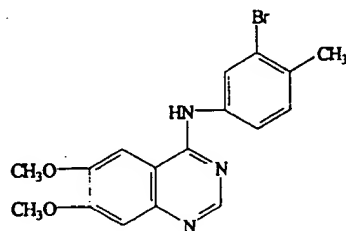
4-(3'-Bromophenyl)-amino-6,7-dimethoxyquinazoline [WHI-P79]



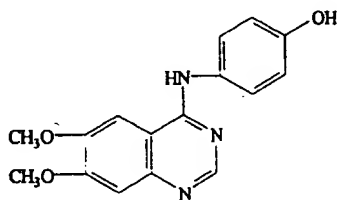
yield 84.17%; m.p. 246.0-249.0°C.
UV(MeOH) λ_{max} : 217.0, 227.0, 252.0 nm;
IR(KBr) ν_{max} : 3409, 2836, 1632, 1512, 1443, 1243,
1068 cm^{-1} ; ^1H NMR(DMSO- d_6): δ 10.42(br, s, 1H,
NH), 8.68(s, 1H, 2-H), 8.07-7.36(m, 5H, 5,
2',4',5',6'-H), 7.24(s, 1H, 8H), 3.98(s, 3H, -
OCH₃), 3.73(s, 3H, -OCH₃); GC/MS m/z 361(M^+
+ 1, 61.83), 360(M^+ , 100), 359(M^+ - 1, 63.52),
344(11.34), 222(10.87), 140(13.65). Anal.
(C₁₆H₁₄BrN₃O₂) C, H, N.

4-(3',5'-Dibromo-4'-hydroxylphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P97]

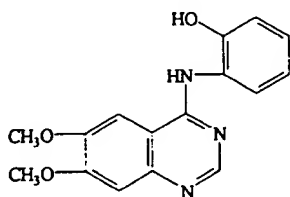
yield 72.80%; m.p. >300.0 °C. UV(MeOH) λ_{max} : 208.0, 210.0, 245.0, 320.0 nm; IR(KBr) ν_{max} : 3504(br), 3419, 2868, 1627, 1512, 1425, 1250, 1155 cm^{-1} ; ^1H NMR(DMSO- d_6): δ 9.71(s, 1H, -NH), 9.39(s, 1H, -OH), 8.48(s, 1H, 2-H), 8.07(s, 2H, 2', 6'-H), 7.76(s, 1H, 5-H), 7.17(s, 1H, 8-H), 3.94(s, 3H, -OCH₃), 3.91(s, 3H, -OCH₃). GC/MS m/z 456(M^+ +1, 54.40), 455(M^+ , 100.00), 454(M^+ -1, 78.01), 439(M^+ -OH, 7.96), 376(M^+ +1 - Br, 9.76), 375(M^+ -Br, 10.91), 360(5.23). Anal. ($\text{C}_{16}\text{H}_{13}\text{Br}_2\text{N}_3\text{O}_3$) C, H, N.

4-(3'-Bromo-4'-methylphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P111]

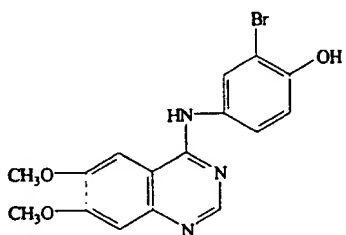
yield 82.22 %; m.p. 225.0-228 °C. ^1H NMR(DMSO- d_6): δ 10.23(s, 1H, -NH), 8.62(s, 1H, 2-H), 8.06(d, 1H, $j_{2,6}=2.1$ Hz, 2'-H), 7.89(s, 1H, 5-H), 7.71(dd, 1H, $J_{5,6}=8.7$ Hz, $J_{2,6}=2.1$ Hz, 6'-H), 7.37(d, 1H, $J_{5,6}=8.7$ Hz, 5'-H), 7.21(s, 1H, 8-H), 3.96(s, 3H, -OCH₃), 3.93(s, -OCH₃). UV(MeOH) λ_{max} (ϵ): 204.0, 228.0, 255.0, 320.0 nm. IR(KBr) ν_{max} : 3431, 3248, 2835, 1633, 1517, 1441, 1281, 1155 cm^{-1} . GC/MS m/z 375(M^+ +1, 76.76), 374(M^+ , 100.00), 373(M^+ -1, 76.91), 358(M^+ +1 - OH, 11.15), 357(1.42), 356(6.31). Anal. ($\text{C}_{17}\text{H}_{16}\text{BrN}_3\text{O}_2\text{HCl}$) C, H, N.

4-(4'-Hydroxyphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P131]

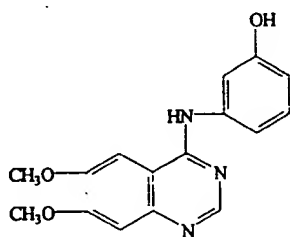
yield 84.29%; m.p. 245.0-248.0 °C. UV(MeOH) λ_{max} : 203.0, 222.0, 251.0, 320.0 nm; IR(KBr) ν_{max} : 3428, 2836, 1635, 1516, 1443, 1234 cm^{-1} ; ^1H NMR(DMSO- d_6): δ 11.21(s, 1H, -NH), 9.70(s, 1H, -OH), 8.74(s, 1H, 2-H), 8.22(s, 1H, 5-H), 7.40(d, 2H, $J = 8.9$ Hz, 2', 6'-H), 7.29(s, 1H, 8-H), 6.85(d, 2H, $J = 8.9$ Hz, 3', 5'-H), 3.98(s, 3H, -OCH₃), 3.97(s, 3H, -OCH₃). GC/MS m/z 298(M^+ +1, 100.00), 297(M^+ , 26.56), 296(M^+ -1, 12.46). Anal. ($\text{C}_{16}\text{H}_{15}\text{N}_3\text{O}_3\text{HCl}$) C, H, N.

4-(2'-Hydroxyphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P132]

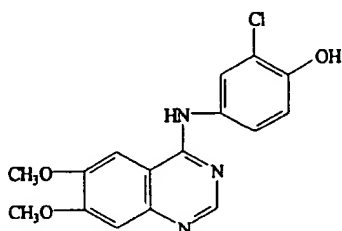
yield 82.49%; m.p. 255.0-258.0 °C. ¹H NMR(DMSO-d₆): δ 9.78(s, 1H, -NH), 9.29(s, 1H, -OH), 8.33(s, 1H, 2-H), 7.85(s, 1H, 5-H), 7.41-6.83(m, 4H, 3', 4', 5', 6'-H), 7.16(s, 1H, 8-H), 3.93(s, 3H, -OCH₃), 3.92(s, 3H, -OCH₃). UV(MeOH)λ_{max}(ε): 203.0, 224.0, 245.0, 335.0 nm. IR(KBr)ν_{max}: 3500 (br), 3425, 2833, 1625, 1512, 1456, 1251, 1068 cm⁻¹. GC/MS m/z 298(M⁺ +1, 8.91), 297(M⁺, 56.64), 281(M⁺ +1- OH, 23.47), 280(M⁺ - OH, 100.00). Anal. (C₁₆H₁₅N₃O₃HC1) C, H, N.

4-(3'-Bromo-4'-hydroxyphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P154]

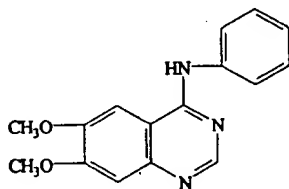
yield 89.90%; m.p. 233.0-233.5 °C. UV(MeOH)λ_{max}: 203.0, 222.0, 250.0, 335.0 nm; IR(KBr) ν_{max}: 3431 br, 2841, 1624, 1498, 1423, 1244 cm⁻¹; ¹H NMR(DMSO-d₆): δ 10.08(s, 1H, -NH), 9.38(s, 1H, -OH), 8.40(s, 1H, 2-H), 7.89(d, 1H, J_{2',5'} = 2.7 Hz, 2'-H), 7.75(s, 1H, 5-H), 7.55(dd, 1H, J_{5',6'} = 9.0 Hz, J_{2',6'} = 2.7 Hz, 6'-H), 7.14(s, 1H, 8-H), 6.97(d, 1H, J_{5',6'} = 9.0 Hz, 5'-H), 3.92(s, 3H, -OCH₃), 3.90(s, 3H, -OCH₃). GC/MS m/z 378(M⁺ +2, 90.68), 377(M⁺ +1, 37.49), 376(M⁺, 100.00), 360(M⁺, 3.63), 298(18.86), 282 (6.65). Anal. (C₁₆H₁₄N₃O₃HCl) C, H, N.

4-(3'-Hydroxyphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P180]

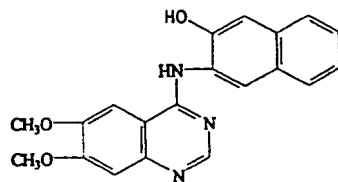
yield 71.55 %; m.p. 256.0- 258.0 °C. ¹H NMR(DMSO-d₆): δ 9.41(s, 1H, -NH), 9.36(s, 1H, -OH), 8.46(s, 1H, 2-H), 7.84(s, 1H, 5-H), 7.84-6.50(m, 4H, 2', 4', 5', 6' -H), 7.20(s, 1H, 8-H), 3.96(s, 3H, -OCH₃), 3.93(s, 3H, -OCH₃). UV(MeOH)λ_{max}(ε): 204.0, 224.0, 252.0, 335.0 nm. IR(KBr)ν_{max}: 3394, 2836, 1626, 1508, 1429, 1251 cm⁻¹. GM/MS m/z: 297(M⁺, 61.89), 296(M⁺, 61.89), 296(M⁺ -1, 100.00), 280(M⁺ - OH, 13.63). Anal. (C₁₆H₁₅N₃O₃.HCl) C, H, N.

4-(3'-Chloro-4'-Hydroxyphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P197]

yield 84.14%; m.p. 245.0°C(dec). ¹H NMR(DMSO-d₆): δ 10.00(s, 1H, -NH), 9.37(s, 1H, -OH), 8.41(s, 1H, 2-H), 7.78(s, 1H, 5-H), 7.49(d, 1H, J_{2,6}=2.7 Hz, 2'-H), 7.55(dd, 1H, J_{5,6}=9.0 Hz, J_{2,6}=2.7 Hz, 6'-H), 7.16(s, 1H, 8-H), 6.97(d, 1H, J_{5,6}=9.0 Hz, 5'-H), 3.93(s, 3H, -OCH₃), 3.91(s, 3H, -OCH₃). UV(MeOH)λ_{max}(ε): 209.0, 224.0, 249.0, 330.0 nm. IR(KBr)ν_{max}: 3448, 2842, 1623, 1506, 1423, 1241 cm⁻¹. GC/MS m/z: 341(M⁺, 100.00), 326(M⁺-CH₃, 98.50), 310(M⁺-OCH₃, 12.5), 295(9.0), 189(13.5), 155(13.8). Anal. (C₁₆H₁₄ClN₃O₃·HCl) C, H, N.

4-(phenyl)-amino-6,7-dimethoxyquinazoline [WHI-P258]

yield 88.6%; m.p. 258.0-260.0 °C. ¹H NMR(DMSO-d₆): δ 11.41(s, 1H, -NH), 8.82(s, 1H, 2-H), 8.32(s, 1H, 5-H), 7.70-7.33(m, 5H, 2', 3', 4', 5', 6'-H), 7.36(s, 1H, 8H), 4.02(s, 3H, -OCH₃), 4.00(s, 3H, -OCH₃), 4.00(s, 3H, -OCH₃). UV(MeOH)λ_{max}(ε): 210.0, 234.0, 330.0 nm. IR(KBr)ν_{max}: 2852, 1627, 1509, 1434, 1248 cm⁻¹. GC/MS m/z 282(M⁺+1, 10.50), 281(M⁺, 55.00), 280(M⁺-1, 100.00), 264(16.00), 207(8.50). Anal. (C₁₆H₁₅N₃O₂) C, H, N.

4-(2'-Hydroxy-naphthyl)-3'- amino-6,7-dimethoxyquinazoline[WHI-P292].

Yield 87.41%; m.p.277.0-279.0°C, IR(KBr)ν_{max}: 3479, 3386, 3036, 2901, 1632, 1581, 1504, 1437, 1281 cm⁻¹. ¹H NMR(DMSO-d₆): δ 11.38(s, 1H, -NH), 10.35(s, 1H, -OH), 8.73(s, 1H, 2-H), 8.25(s, 1H, 5-H), 7.93-7.30(m, 6H, 1', 4', 5', 6', 7', 8'-H), 7.37(s, 1H, 8H), 4.00(s, 6H, -OCH₃). GC/MS m/a: 281(41.0), 253(11.0), 207(100.0). Anal. (C₂₀H₁₇N₃O₃·HCl) C, H, N.

Example 3
Cytotoxicity of Substituted Quinazoline Derivatives

The cytotoxicity of the substituted quinazoline derivative compounds against human glioblastoma cells was evaluated. The relative importance of particular substituent group on the compounds was also studied. The substituted quinazoline derivative compounds, prepared as described above for Example 1, were tested, along with DMSO and Genistein as controls.

10 Cytotoxicity Assay

The cytotoxicity assay of various compounds against human brain tumor cell lines was performed using the MTT (3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyl tetrazolium bromide) assay (Boehringer Mannheim Corp., Indianapolis, IN). Briefly, exponentially growing brain tumor cells were seeded into a 96-well plate at a density of 2.5×10^4 cells/well and incubated for 36 hours at 37°C prior to drug exposure. On the day of treatment, culture medium was carefully aspirated from the wells and replaced with fresh medium containing the quinazoline compounds WHI-P79, WHI-P97, WHI-P111, WHI-P131, WHI-P132, WHI-P154, WHI-P180, WHI-P197, WHI-P258, unconjugated EGF, or EGF-P154, as well as the tyrosine kinase inhibitory isoflavone genistein (GEN) at concentrations ranging from 0.1 to 250 μ M. Triplicate wells were used for each treatment.

Human glioblastoma cell line U373 was obtained from American Type Culture Collection (Rockville, MD) and maintained as a continuous cell line in Dulbecco's modified Eagles's medium supplemented with 10% fetal bovine serum and antibiotics. The B-lineage acute lymphoblastic leukemia cell line Nalm-6 was used as a negative control.

The cells were incubated with the various compounds for 24-36 hours at 37°C in a humidified 5% CO₂ atmosphere. To each well, 10 μ l of MTT (0.5 mg/ml final concentration) was added and the plates were incubated at 37°C for 4 hours to allow MTT to form formazan crystals by reacting with metabolically active cells. The formazan crystals were solubilized overnight at 37°C in a solution containing 10% SDS in 0.01 M HCl. The absorbence of each well was measured in a microplate reader (Labsystems) at 540 nm and a reference wavelength of 690 nm.

To translate the OD₅₄₀ values into the number of live cells in each well, the OD₅₄₀ values were compared to those on standard OD₅₄₀ - versus - cell number curves generated for each cell line. The percent survival was calculated using the formula:

$$\% \text{ Survival} = \frac{\text{live cell number [test]}}{\text{live cell number [control]}} 100$$

The IC₅₀ values were calculated by non-linear regression analysis.

***In situ* Detection of Apoptosis**

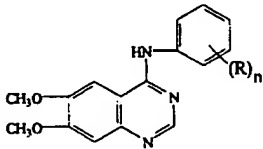
The demonstration of apoptosis was performed by the *in situ* nick-end-labeling method using ApopTag *in situ* detection kit (Oncor, Gaithersburg, MD) according to the manufacturer's recommendations. Exponentially growing cells were seeded in 6-well tissue culture plates at a density of 50x10⁴ cells/well and cultured for 36 hours at 37°C in a humidified 5% CO₂ atmosphere. The supernatant culture medium was carefully aspirated and replaced with fresh medium containing unconjugated EGF or EGF-P154 at a concentration of 10, 25 or 50 µg/ml. After a 36 hour incubation at 37°C in a humidified 5% CO₂ incubator, the supernatants were carefully aspirated and the cells were treated for 1-2 minutes with 0.1% trypsin. The detached cells were collected into a 15 ml centrifuge tube, washed with medium and pelleted by centrifugation at 1000 rpm for 5 minutes. Cells were resuspended in 50 µl of PBS, transferred to poly-L-lysine coated coverslips and allowed to attach for 15 minutes. The cells were washed once with PBS and incubated with equilibration buffer for 10 minutes at room temperature.

After removal of the equilibration buffer, cells were incubated for 1 hour at 37°C with the reaction mixture containing terminal deoxynucleotidyl transferase (TdT) and digoxigenin-11-UTP for labeling of exposed 3'-hydroxyl ends of fragmented nuclear DNA. The cells were washed with PBS and incubated with anti-digoxigenin antibody conjugated to FITC for 1 hour at room temperature to detect the incorporated dUTP. After washing the cells with PBS, the coverslips were mounted onto slides with Vectashield containing propidium iodide (Vector Labs, Burlingame, CA) and viewed with a confocal laser scanning microscope. Non-apoptotic cells do not incorporate significant amounts of dUTP due to lack of

exposed 3-hydroxyl ends, and consequently have much less fluorescence than apoptotic cells which have an abundance of exposed 3'-hydroxyl ends. In control reactions, the TdT enzyme was omitted from the reaction mixture.

5 Results

The identity of the specific substituents on each aniline moiety are summarized below:

	Quinazoline Derivatives	Identity of Substituents
	WHI-P79	3-Br
	WHI-P97	3-Br, 5-Br, 4-OH
	WHI-P111	3-Br, 4-CH ₃
	WHI-P131	4-OH
	WHI-P132	2-OH
	WHI-P154	3-Br, 4-OH
	WHI-P180	3-OH
	WHI-P197	3-Cl, 4-OH
	WHI-258	H
	WHI-292	1-OH Naphthyl

The results of the cytotoxicity assay against brain tumor cells are shown in Figures 1A and 1B. In summary, those compounds having hydroxy substitutions on the phenyl ring, were effective in killing brain tumor cells. The unsubstituted compound P258 and the 3-Br-substituted potent TK inhibitor P79 were ineffective in killing brain tumor cells. Genistein and DMSO controls were also ineffective.

Those substituted quinazoline derivatives having an hydroxyl group on the aniline moiety demonstrated cytotoxic activity. Four compounds tested possessed a single hydroxyl group; at position 4 (WHI-P131), at position 2 (WHI-P132), at position 3 (WHI-P180), and at position 1 (WHI-P292). All four exhibited significant cytotoxicity, with the WHI-P180 (3-OH) compound demonstrating slightly stronger effects than the other two.

Compounds were having both a hydroxyl substituent and a halogen substituents were also potent cytotoxic agents. The structure of WHI-P197 (3-Cl, 4-OH) differs from that of WHI-P131 (4-OH) only in the chlorine atom at position 3. As shown in Figures 1A and 1B, addition of the chlorine atom did not effect the cytotoxicity of this compound.

The structure of WHI-P154 (3-Br, 4-OH) differs from WHI-P131 (4-OH), only in the bromine atom at position 3. As shown in Figures 1A and 1B,

the addition of the bromine atom to this compound significantly increased the cytotoxicity of the compound.

The structure of WHI-P97 differs from that of WHI-P154 only in the additional bromine atom added at position 5. Figure 1 shows that there is essentially
5 no benefit from the added second bromine atom.

WHI-P154, 4-(3'-Bromo-4'-hydroxyphenyl)-amino-6,7-dimethoxyquinazoline exhibited significant cytotoxicity against the U373 human glioblastoma cell line in 3 of 3 independent experiments with a mean (\pm SE) IC₅₀ value of 167.4 ± 26.9 μ M and a composite survival curve IC₅₀ value of 158.5 μ M. In
10 contrast, WHI-P79, a potent inhibitor of EGF-R and Src family tyrosine kinases (Bos, et al., 1997, *Clin. Cancer Res.* 3:2099-2106; Fry, et al., 1994, *Science (Washington, D.C.)* 265:1093-1095) failed to cause any detectable cytotoxicity to U373 glioblastoma cells. Thus, the cytotoxicity of WHI-P154 to U373 cells cannot be explained by its tyrosine kinase inhibitory properties. This notion was further
15 supported by the inability of the PTK inhibitor genistein (included as controls) to cause detectable cytotoxicity to U373 cells (IC₅₀ value >250 μ M; Figure 1A).

Example 4

Enhanced Cytotoxicity of Conjugated WHI-P154

Against Human Glioblastoma Cells

In contrast to normal glial cells and neurons, significant numbers of glioblastoma cells express the EGF receptor (EGF-R) at high levels. Therefore, the EGF-R is a potential target for delivering cytotoxic agents to glioblastoma cells
25 with greater efficiency (Mendelsohn, J. and Baselga, J., 1995, *Biologic Therapy of Cancer: Principles and Practice*, pp. 607-23).

Expression of EGF-R by Glioblastoma Cells

Surface expression of the EGF-R on the U373 and U87 human
30 glioblastoma cell lines (obtained and maintained as described for Example 3) was confirmed with immunofluorescence and confocal laser scanning microscopy using monoclonal antibodies to the extracellular domain of the EGF-R.

Immunofluorescence staining with anti- α -tubulin antibody and the nuclear dye Toto-3 was used in combination with confocal laser scanning microscopy to
35 examine the morphological features of U373 glioma cells treated with either

unconjugated EGF or EGF-P154. Cells were fixed in paraformaldehyde, immunostained with monoclonal antibody to EGF-R (green fluorescence) and counterstained with TOTO-3 (blue fluorescence). The immunostained cells were analyzed with a laser scanning confocal microscope. Blue fluorescence represents nuclei. Both cell lines showed a diffuse granular immunoreactivity with the anti-EGF-R antibody (see Figures 2A and 2B).

Preparation of EGF-P154 Conjugate

In an attempt to enhance the demonstrated anti-tumor activity of 4-(3'-Bromo-4'-hydroxyphenyl)-amino-6,7-dimethoxyquinazoline (WHI-P154) against glioblastoma cells, by improving its targeting to and cellular uptake by glioblastoma cells, the compound was conjugated to recombinant human EGF, as described below.

Recombinant human EGF (rhEGF) was produced in *E. coli* harboring a genetically engineered plasmid that contains a synthetic gene for human EGF fused at the N-terminus to a hexapeptide leader sequence for optimal protein expression and folding. The rhEGF fusion protein precipitated in the form of inclusion bodies and the mature protein was recovered by trypsin-cleavage followed by purification using ion exchange chromatography and HPLC. The recovered rhEGF was 99% pure by reverse-phase HPLC and SDS-PAGE with an isoelectric point of 4.6 ± 0.2 . The endotoxin level was 0.172 EU/mg.

The recently published photochemical conjugation method using the hetero-bifunctional photoreactive crosslinking agent, Sulfosuccinimidyl 6-[4'-azido-2'-nitrophenylamino] hexanoate (Sulfo-SANPAH) (Pierce Chemical Co., Rockford, IL) was employed for the synthesis of the EGF-P154 conjugate, as described in Uckun, et al., 1995, *Science*, 267:886-891. Sulfo-SANPAH modified rhEGF was mixed with a 10:1 molar ratio of WHI-P154 [50 mM solution in dimethyl sulfoxide (DMSO)] and then irradiated with gentle mixing for 10 minutes with UV light at wavelengths 254-366 nm with a multiband UV light-emitter (Model UVGL-15 Mineralight; UVP, San Gabriel, CA). Photolytic generation of a reactive singlet nitrene on the other terminus of EGF-SANPAH in the presence of a 10-fold molar excess of WHI-P154 resulted in the attachment of WHI-P154 to EGF.

Excess WHI-P154 in the reaction mixture was removed by passage through a pre-packed PD-10 column, and 12 kDa EGF-EGF homoconjugates with or without conjugated WHI-P154 as well as higher molecular weight reaction products were removed by size-exclusion high-performance liquid chromatography (HPLC).

- 5 Reverse phase HPLC using a Hewlett-Packard (HP) 1100 series HPLC instrument was used for separation of EGF-P154 from EGF-SANPAH. After the final purification, analytical HPLC was performed using a Spherisorb ODS-2 reverse phase column (250x4 mm, Hewlett-Packard, Cat.# 79992402-584). Prior to the HPLC runs, a Beckman DU 7400 spectrophotometer was used to generate a UV
10 spectrum for each of the samples to ascertain the λ_{max} for EGF-P154, EGF-SANPAH, and unmodified EGF.

- Each HPLC chromatogram was subsequently run at wavelengths of 214, 265, and 480 nm using the multiple wavelength detector option supplied with the instrument to ensure optimal detection of the individual peaks in the
15 chromatogram. Analysis was achieved using a gradient flow consisting of 0% to 100% eluent in a time interval of 0 to 30 minutes. Five μL samples applied to the above column were run using the following gradient program: 0-5 minutes: 0-20% eluent; 5-20 minutes: 20-100% eluent; 25-30 minutes: 100% eluent; and 30-35 minutes: 100-0% eluent. The eluent was a mixture of 80% acetonitrile (CH_3CN),
20 20% H_2O and 0.1% TFA. Electrospray ionization mass spectrometry (Feng, et al., 1991, *J. Am. Soc. Mass Spectrometry* 2:387-401; Covey, et al., 1988, *Rapid Communications in Mass Spectrometry* 2:249-256) was performed using a PE SCIEX API triple quadrupole mass spectrometer (Norwalk, CT) to determine the stoichiometry of P154 and EGF in EGF-P154.

25

Uptake and Internalization of EGF-P154

- The kinetics of uptake and cytotoxicity of the EGF-P154 conjugate in U373 glioblastoma cells were analyzed using immunofluorescence and confocal laser microscopy for following the internalized EGF-R and EGF-P154 molecules,
30 as well as morphologic changes in treated cells.

Immunofluorescence was used to (i) examine the surface expression of EGF-receptor (EGF-R) on brain tumor cells, (ii) evaluate the uptake of EGF-

P154 by brain tumor cells and (iii) examine the morphologic features of EGF-P154 treated brain tumor cells. For analysis of EGF-R expression and cellular uptake of EGF-P154, U87 and U373 glioblastoma cells were plated on poly-L-lysine coated glass-bottom 35 mm Petri dishes (Mattek Corp., Ashland, MA) and
5 maintained for 48 hours. In uptake studies, the culture medium was replaced with fresh medium containing 5 µg/ml EGF, EGF-P154 or unconjugated WHI-P154 and cells were incubated at 37°C for 5 minutes, 10 minutes, 15 minutes, 30 minutes, 60 minutes, and 24 hours.

At the end of the incubation, cells were washed twice with PBS and
10 fixed in 2% paraformaldehyde. The cells were permeabilized and non-specific binding sites were blocked with 2.5% BSA in PBS containing 0.1% Triton X-100 for 30 minutes. To detect the EGF-R/EGF-P154 complexes, cells were incubated with a mixture of a monoclonal antibody (1:10 dilution in PBS containing BSA and Triton X-100) directed to the extracellular domain of the human EGF-R (Santa
15 Cruz Biotechnologies Inc., Santa Cruz, CA) and a polyclonal rabbit anti-P154 antibody (1:500 dilution) for 1 hour at room temperature. After rinsing with PBS, cells were incubated for 1 hour with a mixture of a goat anti-mouse IgG antibody conjugated to FITC (Amersham Corp., Arlington Heights, IL) and donkey anti-rabbit IgG conjugated to Texas Red (Amersham Corp.) at a dilution of 1:40 in
20 PBS. Similarly, tubulin expression was examined by immunofluorescence using a monoclonal antibody against α -tubulin (Sigma Chemical Co, St. Louis, MO) at a dilution of 1:1000 and an anti-mouse IgG conjugated to FITC. Cells were washed in PBS and counterstained with toto-3 (Molecular Probes Inc., Eugene, OR) for 10 minutes at a dilution of 1:1000. Cells were washed again with PBS and the
25 coverslips were mounted with Vectashield (Vector Labs, Burlingame, CA) and viewed with a confocal microscope (Bio-Rad MRC 1024) mounted in a Nikon Labhophot upright microscope.

Cytotoxic Activity

30 To determine if the improved delivery of WHI-P154 to glioblastoma cells by conjugation to EGF resulted in potentiation of its anti-tumor activity, the cytotoxic activities of EGF-P154 and unconjugated WHI-P154

against U373 and U87 human glioblastoma cell lines were analyzed in dose response studies using *in vitro* MTT assays, as described above for Example 3. Apoptosis was also evaluated using the nick-end-labeling assay, as described for Example 3. The data are discussed below.

5 Results

Surface expression of the EGF-R on the U373 and U87 human glioblastoma cell lines was confirmed with immunofluorescence and confocal laser scanning microscopy using monoclonal antibodies to the extracellular domain of the EGF-R. As shown in Figures 2a and 2b, both cell lines showed a diffuse granular
10 immunoreactivity with the anti-EGF-R antibody.

The kinetics of uptake and cytotoxicity of the EGF-P154 conjugate in U373 glioblastoma cells were analyzed using immunofluorescence and confocal laser microscopy for following the internalized EGF-R and EGF-P154 molecules, as well as morphologic changes in treated cells. EGF-P154, similar to unconjugated
15 EGF (not shown), was able to bind to and enter target glioblastoma cells via receptor-mediated endocytosis by inducing internalization of the EGF-R molecules. As shown in Figures 3A-C, 3C', 3D and 3D', within 10 minutes after exposure to EGF-P154, the EGF-R/EGF-P154 complexes began being internalized, as determined by co-localization of the EGF-R (detected by anti-EGF-R antibody,
20 green fluorescence) and EGF-P154 (detected by anti-P154 antibody, red fluorescence) in the cytoplasm of treated cells. By 30 minutes, the internalized EGF-R/EGF-P154 complexes were detected in the perinuclear region of the treated glioblastoma cells. In contrast, cells treated with unconjugated WHI-P154 alone (Figures 3B, 3B') did not reveal any detectable redistribution of the surface EGF-R
25 or cytoplasmic staining with the anti-P154 antibody (red fluorescence). By 24 hours (but not at 6 or 12 hours), WHI-154 molecules could also be detected in cells treated with unconjugated WHI-154 (Figures 4A, 4A', 4B, 4B'). Thus, conjugation of WHI-P154 to EGF resulted in increased uptake of this cytotoxic quinazoline derivative by EGF-R positive glioblastoma cells.

30 To determine if the improved delivery of WHI-P154 to glioblastoma cells by conjugation to EGF resulted in potentiation of its anti-tumor activity, the cytotoxic activities of EGF-P154 and unconjugated WHI-P154 against U373 and U87 human glioblastoma cell lines were analyzed in dose response studies using *in vitro* MTT assays, as described above for Example 3. As shown in Figure 5A,
35 EGF-P154 killed these glioblastoma cells in each of 3 independent experiments at

nanomolar concentrations with mean IC₅₀ values of 813 ± 139 nM (Range: 588 – 950 nM) for U373 cells and 620 ± 97 nM (Range: 487 – 761 nM) for U87 cells. The IC₅₀ values derived from the composite cell survival curves were 881 nM for U373 cells and 601 nM for U87 cells. By comparison, unconjugated WHI-P154 killed

5 U373 or U87 cells only at micromolar concentrations. EGF-P154 was 206-fold more potent than unconjugated WHI-P154 against U373 cells (IC₅₀ values: 167.4 ± 26.9 μ M vs 811 ± 139 nM, $P < 0.003$) and 288-fold more potent than unconjugated WHI-P154 against U87 cells (IC₅₀ values: 178.62 ± 18.46 μ M vs 620 ± 97 nM, $P < 0.001$) (**Figure 5A**).

10 Unlike WHI-P154, which showed marked cytotoxicity against the EGF-R negative NALM-6 leukemia cells, EGF-P154 elicited selective cytotoxicity to EGF-R positive glioblastoma cell lines only (**Figures 5A and 5B**). Thus, conjugation to EGF increased the potency of WHI-P154 against human glioblastoma and at the same time restricted its cytotoxicity to EGF-R positive

15 targets.

Unlike the EGF-P154 conjugate, EGF-GEN, a potent inhibitor of the EGF-R tyrosine kinase and EGF-R associated Src family PTK, failed to kill glioblastoma cells (**Figure 5A**). Thus, the potent cytotoxicity of EGF-P154 cannot be explained by the tyrosine kinase inhibitory properties of its WHI-P154 moiety.

20 Immunofluorescence staining with anti- α -tubulin antibody and the nuclear dye TOTO-3 was used in combination with confocal laser scanning microscopy to examine the morphological features of U373 glioma cells treated with either unconjugated EGF or EGF-P154. As shown in **Figures 6A–6C**, after 24 hours of exposure to 25 μ g/ml EGF-P154 (but not 25 μ g/ml unconjugated EGF),

25 most of the glioma cells showed an abnormal architecture with complete disruption of microtubules, marked shrinkage, nuclear fragmentation and inability to adhere to the substratum. These morphologic changes in EGF-P154-treated glioma cells were consistent with apoptosis.

To confirm apoptotic DNA fragmentation in the nuclei of EGF-P154-treated glioblastoma cells, an *in situ* apoptosis assay which allows the

30 detection of exposed 3'-hydroxyl groups in fragmented DNA by TdT-mediated dUTP nick-end labeling was used. As evidenced by the confocal laser scanning microscopy images depicted in **Figures 6D and 6E**, EGF-P154 treated (but not EGF-treated) glioma cells examined for digoxigenin-dUTP incorporation using

35 FITC-conjugated anti-digoxigenin (green fluorescence) and propidium iodide

counterstaining (red fluorescence) showed many apoptotic yellow nuclei with superimposed green and red fluorescence at 36 hours after treatment.

In summary, these data demonstrate specific and enhanced cytotoxic activity of P154 when targeted to brain tumor cells by conjugation to EGF.

5

Example 5

Anti-tumor Activity of EGF-P154 in a SCID Mouse Model

The anti-tumor effects of conjugated EGF-P154 *in vivo* is demonstrated in a SCID mouse model. CB.17 SCID mice developed rapidly growing tumors after subcutaneous inoculation of 0.5×10^6 U373 cells. The *in vivo* anti-tumor activity of EGF-P154 in this SCID mouse xenograft model of human glioblastoma multiforme was examined.

10

Maintenance of SCID Mouse Colony

15

The SCID mice were housed in a specific pathogen-free room located in a secure indoor facility with controlled temperature, humidity, and noise levels. The SCID mice were housed in microisolator cages which were autoclaved with rodent chow. Water was also autoclaved and supplemented with trimethoprim/sulfamethoxazole 3 days/week.

20

SCID Mouse Xenograft Model of Human Glioblastoma

The right hind legs of the CB. 17 SCID mice were inoculated subcutaneously (s.c.) with 0.5×10^6 U373 human glioblastoma cells in 0.2 mL PBS. SCID mice challenged with brain tumor cells were treated with EGF-P154 (0.5 mg/kg/dose or 1 mg/kg/dose in 0.2 ml PBS) as daily i.p. doses for 10 treatment days starting the day after inoculation of the glioblastoma cells. Daily treatments with PBS, unconjugated EGF (1 mg/kg/dose), and unconjugated WHI-P154 (1 mg/kg/dose) were used as controls. Mice were monitored daily for health status and tumor growth, and were sacrificed if they became moribund, developed tumors which impeded their ability to attain food or water, at the end of the 3-month observation period. Tumors were measured using Vernier calipers twice weekly, and the tumor volumes were calculated according to the following formula

25

30

(Friedman, S.H., mDolan, M.E., Pegg, A.E., Marcelli, S., Keir, S., Catino, J.J., Binger, D.D., Schold, S.C., Jr., 1995, *Cancer Res.*, 55:2853-2857):

$$\text{Volume} = \frac{\text{Width}^2 * \text{Length}}{2}$$

5

For histopathologic studies, tissues were fixed in 10% neutral buffered formalin, dehydrated, and embedded in paraffin by routine methods. Glass slides with affixed 6 micron tissue sections were prepared and stained with hematoxylin/eosin. Primary endpoints of interest were tumor growth and tumor - free survival outcome. Estimation of life table outcome and comparisons of outcome between groups were done, as previously reported (Waurzyniak, et al., 1997, *Clinical Cancer Research* 3:881-890; Anderson, et al., 1995, *Cancer Res.* 55:1321-1327; Uckun, et al., 1997, *J. Clin. Oncol.* 15:2214-2221).

15 Results

The conjugated quinazoline EGF-P154 significantly improved tumor-free survival in a dose-dependent fashion, when it was administered 24 hours after inoculation of tumor cells. **Figures 7A and 7B** show the tumor growth and tumor-free survival outcome of SCID mice treated with EGF-P154 (500 µg/kg/day x 10 days or 1 mg/kg/day x 10 days), unconjugated EGF (1 mg/kg/day x 10 days), unconjugated WHI-P154 (1 mg/kg/day x 10 days), or PBS after inoculation with U373 glioblastoma cells.

None of the 15 control mice treated with PBS (N= 5; median tumor-free survival = 19 days), EGF (N= 5; median tumor-free survival = 23 days), or unconjugated WHI-P154 (N= 5; median tumor-free survival = 19 days) remained alive tumor-free beyond 33 days (median tumor-free survival = 19 days) (**Figure 7A**). All of the 5 mice treated with EGF-P154 at the 500 µg/kg/day dose level developed tumors within 40 days with an improved median tumor-free survival of 33 days (**Figure 7B**). These tumors were much smaller than those in control mice (**Figure 7A**).

30

Tumors reached a size of 50 mm³ by 37.5±3.3 days in PBS treated mice, 34.0±3.0 days in EGF-treated mice, and 36.0±5.1 days in WHI-P154 treated mice. Tumors developing in EGF-P154 (500 µg/kg/day x 10 days)-treated mice reached the 50 mm³ tumor size approximately 11 days later than the tumors in

control mice treated with PBS, EGF, or WHI-P154 (47.4 ± 7.1 days vs 35.8 ± 1.8 days). The average size (mean \pm SE) of tumors at 20 days and 40 days were 10.2 ± 1.4 mm³ and 92.3 ± 6.0 mm³, respectively for mice in the control groups (i.e., PBS + EGF groups combined). By comparison, the average size (mean \pm SE) of tumors at
5 20 days and 40 days were significantly smaller at 1.0 ± 1.1 mm³ ($P = 0.002$) and 37.6 ± 10.7 mm³ ($P = 0.0003$) for mice treated with EGF-P154 at the 500 μ g/kg/day dose level.

Notably, 40 % of mice treated for 10 consecutive days with 1 mg/kg/day EGF-P154 remained alive and free of detectable tumors for >58 days
10 (PBS + EGF + WHI-P154 vs EGF-P154, $P < 0.00001$ by log-rank test). The tumors developing in the remaining 60% of the mice did not reach a size >50 mm³ during the 58-day observation period. Thus, EGF-P154 elicited significant *in vivo* anti-tumor activity at the applied nontoxic dose levels. The inability of 1 mg/kg/day x 10
15 days of unconjugated WHI-P154 (53.2 nmol) and unconjugated EGF to confer tumor-free survival in this SCID mouse model in contrast to the potency of 1 mg/kg/day x 10 days EGF-P154 (corresponding to 2.9 nmol of WHI-P154) demonstrates that the *in vivo* anti-tumor activity of EGF-P154 cannot be attributed to its EGF moiety alone and also that conjugation to EGF enhances the *in vivo* anti-tumor activity of WHI-P154 against glioblastoma cells by >18 -fold.

20 Taken together, the findings of Examples 3-5 provide unprecedented evidence that the substituted quinazoline 3-bromo-4'-hydroxyphenyl)-amino-6,7-dimethoxyquinazoline (WHI-P154) exhibits significant cytotoxicity against human glioblastoma cells and that its anti-tumor activity can be substantially enhanced by conjugation to EGF as a targeting molecule. Although WHI-P154 is a
25 potent inhibitor of the EGF-R kinase as well as Src family tyrosine kinases, its cytotoxicity in glioblastoma cells cannot be attributed to its tyrosine kinase inhibitory properties alone, since 4-(3'-Bromophenyl)-amino-6,7-dimethoxyquinazoline (WHI-P79) with equally potent PTK inhibitory activity, failed to kill WHI-P154 sensitive glioblastoma cells. Similarly, several PTK
30 inhibitors capable of killing human leukemia and breast cancer cells lacked detectable cytotoxicity against glioblastoma cells. Glioblastoma cells exposed to EGF-conjugated WHI-P154 underwent apoptosis. Although EGF was used to target WHI-P154 to glioblastoma cells in the present study, other biologic agents including different cytokines such as IGF and antibodies reactive with glioblastoma-

associated antigens are also expected to be effective targeting molecules for this novel quinazoline derivative.

Example 6

5 Substituted Quinazolines Inhibit Glioblastoma Cell Adhesion

During the multistep process of tissue invasion, tumor cells initially adhere to the extracellular matrix proteins via cell surface integrin receptors and then gain migratory capacity to enter the surrounding tissues. ECM proteins such as laminin, fibronectin, and type IV collagen are thought to play an important role in
10 tumor cell attachment and migration. Laminin, fibronectin and collagen have been found in the basal lamina of blood vessels and in the glial limitans externa in the brain that promote the adhesion and invasion of tumor cells *in situ* (Carbonetto, S., 1984, *Trends Neurosci.*, 7:382-387; Rutka, J.T., Apodaca, G., Stern, R., *J. Neurosurg.*, 69:155-170; Venstrom, K.A. and Reichard, L.F., 1993, *FASEB J.*,
15 7:996-1003). The effects of these ECM proteins on integrin-mediated glioblastoma cell adhesion was examined using four different human glioblastoma (U87, U373, T98, and U138) and one medulloblastoma (Daoy) cell line.

Cell Lines

20 Human brain tumor cell lines derived from adult patients with glioblastoma, U-87 MG (Cat.#HTB-14), U-118 MG (Cat. #HTB-15), U-138 MG (Cat. #HTB 16), U-373 MG (Cat. #HTB-17), T98-G (Cat. #CRL-1690) and medulloblastoma Daoy (Cat. #HTB-186) were obtained from American Type Culture Collection (ATCC, Rockville, MD) and maintained in liquid culture using
25 DMEM supplemented with 10% fetal bovine serum and antibiotics. Fibroblast conditioned medium was used as a source of chemoattractant in *in vitro* invasion assays. Conditioned medium was prepared as described previously (Albini, A., Iwamoto, Y., Kleinman, H.K., Martin, G.R., Aaronson, S.A., Kozlowski, J.M., and McEwan, R.N., 1987, *Cancer Res.*, 47:3239-3245). For the preparation of this
30 conditioned medium NIH/3T3 embryonic fibroblasts (AATCC cat. #CRL-1658) were grown to 80% confluency in DMEM medium supplemented with 10% FBS and cultured for 24 hours in serum-free medium containing 0.5 µg/ml bovine serum

albuminates The culture supernatants were collected, centrifuged at 1000 x g for 15 minutes to remove cellular debris and used as conditioned medium.

Adhesion Assays

5 *In vitro* adhesion assays were performed to (a) study the baseline adhesive properties of various glioblastoma cell lines and (b) evaluate the effects of quinazoline derivatives on the adhesive properties of glioblastoma cells. The plates for the adhesion assays were precoated with the extracellular matrix proteins laminin, fibronectin or type IV collagen (each at a final concentration of 1 µg/ml in
10 PBS) overnight at 4°C and dried. On the day of the experiment, the wells were rehydrated and blocked with 10% bovine serum albumin in PBS for 1 hour at room temperature and used for the adhesion assays, as described below.

To study the effects of quinazoline derivatives on glioblastoma cell adhesion, exponentially growing cells in DMEM were incubated with the
15 compounds WHI-P79, WHI-P97, WHI-P31, WHI-P154, WHI-P258 or genistein at concentrations ranging from 1 µM to 100 µM for 16 hours in a humidified 5% CO₂ atmosphere. DMSO (0.1%) was included as a vehicle control. After treatment, cells were detached from the flasks with 0.05% trypsin (Life Technologies) resuspended in DMEM, incubated at 37°C for 2 hours to allow them to recover from the
20 trypsinization stress and examined for their ability to adhere to plates precoated with ECM proteins.

In adhesion assays, cells were centrifuged, washed twice with serum-free DMEM, counted and resuspended in serum-free DMEM to a final concentration of 2.5×10^5 cells/ml. One hundred µl of the cell suspension containing 2.5×10^4 cells
25 were added to each well and cells were allowed to adhere for 1 hour at 37°C in a humidified 5% CO₂ atmosphere. The adherent fraction was quantitated using MTT (3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide) assays. In brief, after washing the wells, 10 µl of MTT (0.5 mg/ml final concentration) (Boehringer Mannheim Corp., Indianapolis, IN) was added to each well and the plates were
30 incubated at 37°C for 4 hours to allow MTT to form formazan crystals by reacting with metabolically active cells. The formazan crystals were solubilized overnight at 37°C in a solution containing 10% SDS in 0.01 M HCl. The absorbance of each

well was measured in a microplate reader (Labsystems) at 540 nm and a reference wavelength of 690 nm. To translate the OD₅₄₀ values into the number of cells in each well, the OD₅₄₀ values were compared to those on standard OD₅₄₀-versus-cell number curves generated for each cell line. The adherent fractions of cells treated with quinazoline derivatives were compared to those of DMSO-treated control cells and the percent inhibition of adhesion was determined using the formula:

$$\% \text{ Inhibition} = 100 * \frac{1 - \text{Adherent Fraction of Drug Treated Cells}}{\text{Adherent Fraction of Control Cells}}$$

- Each treatment condition was evaluated in duplicate in 3 independent experiments. The IC₅₀ values were calculated by non-linear regression analysis.

Results

As shown in **Figure 8**, a significantly greater fraction of glioblastoma and medulloblastoma cells adhered to plates precoated with laminin, type IV collagen, or fibronectin than to uncoated or poly-L-lysine-coated control plates. Of the four glioblastoma cell lines examined, U373 cells were the most adhesive. Therefore, U373 cells were used in subsequent experiments that were designed to examine the effects of various quinazoline derivatives on integrin-mediated glioblastoma cell adhesion.

As shown in **Figures 9A–9D**, the novel quinazoline derivative 4-(3'-Bromo-4'-hydroxyphenyl)-amino-6,7-dimethoxyquinazoline (WHI-P154) (but not the unsubstituted parent compound WHI-P258) inhibited the adhesion of U373 cells to laminin-, fibronectin-, and collagen-coated plates in a dose-dependent fashion with mean IC₅₀ values of 29.8±3.1 μM (N=3) for adhesion to fibronectin-coated plates, 36.1±3.5 μM (N=3) for adhesion to laminin-coated plates, and 42.7±2.5 μM (N=3) for adhesion to collagen-coated plates. The 3'-bromo substitution on the phenyl ring likely contributes to the activity of WHI-P154 since 4-(4'-hydroxyphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P131] lacking this bromo substituent was less potent than WHI-P154 (all IC₅₀ values: >50 μM). The 4'-hydroxyl substituent on the phenyl ring also contributed to the inhibitory activity of WHI-P154 since 4-(3'-Bromophenyl)-amino-6,7-dimethoxyquinazoline

[WHI-P79] which differs from WHI-P154 only by the lack of the 4'-hydroxyl group on the phenyl ring, was less potent (all IC₅₀ values: >50 μ M). Introduction of a second bromo group at the 5' position of the phenyl ring did not result in improved inhibitory activity: 4-(3',5'-Dibromo-4'-hydroxylphenyl)-amino-6,7-dimethoxyquinazoline [WHI-P97] was not more potent than WHI-P154.

Example 7

Effect of WHI-P154 on EGF-induced Cell Adhesion

In addition to binding of cell surface receptors to ECM proteins and integrin clustering, formation of focal adhesion plaques is also regulated by the activation of focal adhesion kinase by certain growth factors upon binding to their receptors (Hatai, M., Hashi, H., Mogi, A., Soga, H., Yokota, J., Yaoi, Y., 1994, *FEBS Lett.*, 350:113-116; Ouwens, D.M., Mikkers, H.M., van der Zon, G.C., Stein Gerlach, M., Ullrich, A., Maassen, J.A., 1996, *Biochem J.*, 318:609-614; Schaller, M.D., Parsons, J.T., *Curr. Opin. Cell*; Zachary, I., 1997, *Int. J. Biochem., Cell Biol.*, 29:929-934). EGF is a potent mitogen for various brain tumor cells that express EGF receptor and are also shown to modulate the expression of cell surface

To study the effects of quinazoline derivatives on EGF-stimulated cell adhesion, the trypsinized and recovered cells were incubated with varying concentrations ranging from 1 μ M to 50 μ M of quinazolines for 4 hours at 37°C, then stimulated with 250 ng/ml of EGF and examined for their ability to adhere to poly-L-lysine coated plates.

For the EGF stimulation experiments, the cells were plated in the presence of 250 ng/ml of EGF and allowed to adhere for 1 hour. The non-adherent cells were removed by gently washing the cells with PBS and then the adherent fraction was quantitated as described above for Example 6.

Stimulation of U373 cells with EGF significantly increased the fraction capable of adhering to poly-L-lysine coated plates from $33.2 \pm 5.2\%$ to $58.48 \pm 4.7\%$ ($P < 0.02$). A four-hour pretreatment of U373 cells with WHI-P154 not only completely prevented the EGF-induced adhesion but it also reduced the adhesive fraction of U373 cells in a dose-dependent fashion far below the baseline levels despite the presence of EGF (Figure 10). Similar, albeit less potent, inhibitory effects were observed when cells were pretreated with WHI-P131.

Example 8

Substituted Quinazolines Inhibit Glioblastoma Cell Invasion

Glioblastoma Cell Invasion Through Matrigel Matrix

5 The *in vitro* invasiveness of glioblastoma cells was assayed using a previously published method which employs Matrigel-coated Costar 24-well transwell cell culture chambers ("Boyden chambers") with 8.0- μ m-pore polycarbonate filter inserts (Albini, A., Iwamoto, Y., Kleinman, H.K., Martin, G.R., Aaronson, S.A., Kozlowski, J.M., and McEwan, R.N., 1987, *Cancer Res.*, 47:3239-
10 3245). The chamber filters were coated with 50 μ g/ml of Matrigel matrix, incubated overnight at room temperature under a laminar flow hood and stored at 4°C. Matrigel matrix is made up of several components of the extracellular matrix (ECM), including collagens, laminin and proteo-glycans

On the day of the experiment, the coated inserts were rehydrated with
15 0.5 ml serum-free DMEM containing 0.1% bovine serum albumin for 1-2 hours. To study the effects of quinazoline derivatives on invasiveness of glioblastoma cells, exponentially growing cells were incubated overnight with WHI-P97, WHI-P131 and WHI-P154 at various concentrations ranging from 1 μ M to 50 μ M. The cells were trypsinized, washed twice with serum-free DMEM containing BSA, counted
20 and resuspended at 1×10^5 cells/ml. 0.5 ml cell suspension containing 5×10^4 cells in a serum-free DMEM containing quinazoline compounds or vehicle was added to the Matrigel-coated and rehydrated filter inserts. Next, 750 μ l of NIH fibroblast conditioned medium was placed as a chemoattractant in 24-well plates and the inserts were placed in wells and incubated at 37°C for 48 hours. After the
25 incubation period, the filter inserts were removed, the medium was decanted off and the cells on the top side of the filter that did not migrate were scraped off with a cotton-tipped applicator. The invasive cells that migrated to the lower side of the filter were fixed, stained with Hema-3 solutions and counted under microscope. Five to 10 random fields per filter were counted to determine the mean (\pm SE) values
30 for the invasive fraction. The invasive fractions of cells treated with quinazoline derivatives were compared to those of DMSO treated control cells and the percent inhibition of invasiveness was determined using the formula:

$$\% \text{ Inhibition} = 100 \left[\frac{1 - \text{Adherent Fraction of Drug Treated Cells}}{\text{Adherent Fraction of Control Cells}} \right]$$

Each treatment condition was evaluated in duplicate in 3 independent experiments.

- 5 IC50 values were calculated by non-linear regression analysis using Graphpad Prism Software Version 2.0 (Graphpad Software Inc., San Diego, CA).

Results

- As shown in **Figure 11**, U373 glioblastoma cells were highly
 10 invasive in Matrigel-coated Boyden chambers. WHI-P154 inhibited the invasion of U373 cells through the Matrigel matrix in dose-dependent fashion and it was more potent than WHI-P131 or WHI-P97 (**Figure 11**). The mean IC50 values obtained from 3 independent experiments were $10.59 \pm 1.8 \mu\text{M}$ (range: $9.57\text{--}11.64 \mu\text{M}$) for WHI-P97, $7.07 \pm 1.8 \mu\text{M}$ (range $5.08\text{--}8.59 \mu\text{M}$) for WHI-P131, and $4.46 \pm 0.8 \mu\text{M}$
 15 (range: $3.53\text{--}5.01$) for WHI-P154. The IC50 values derived from the average values of three experiments were $9.58 \mu\text{M}$ for WHI-P97, $7.95 \mu\text{M}$ for WHI-P131 and $5.2 \mu\text{M}$ for WHI-P154.

Example 9

- 20 **Substituted Quinazolines Inhibit Focal Adhesion Plaques
and Actin Polymerization**

- Cytoskeletal organization and cellular adhesion are two crucial determinants of cell motility and these processes are controlled by the complex coordination of actin cytoskeletal rearrangement and changes in focal adhesions
 25 (Symons, M.H., and Mitchison, T.J., 1991, *J. Cell Biol.*, 114:503–513; Wang, Y.L., 1984, *J. Cell Biol.*, 99:1478–1485; Bretcher, M.S., 1996, *Cell*, 87:601–606; Machesky, L.M., and Hall, A., 1997, *J. Cell Biol.*, 138:913–926). Polymerization of actin filaments, formation of lamellepodia and filapodia at the leading edges are essential for the attachment and detachment of cells from the ECM and play pivotal
 30 roles in cell motility and migration (Burrige, K., Fath, K., Kelly, G., and Turner, C., 1988, *Ann. Rev. Cell Biol.*, 4:487–525; Burrige, K., Nuckolls, C., Otey, F., Pavalko, K., Simon, K., and TAurner, C., 1990, *Cell Differ. Dev.*, 32:337–342). Formation of adhesion plaques is also important in this process because the polymerized actin

fibers are tethered and linked to ECM at these junctions and the cell movement is dependent on the strength of these focal adhesions. Moderate level of cellular adhesive strength is thought to be necessary for cell migration (Burridge, K., Fath, K., Kelly, G., and Turner, C., 1988, *Ann. Rev. Cell Biol.*, 4:487-525; Burridge, K., and Fath, K., 1989, *Bioessays*, 10:104-108; Schwarzbauer, J.E., 1997, *Curr. Biol.*, 7:292-294). Adhesions that are too strong may impair cell motility and adhesions that are too weak may not provide sufficient momentum to move the cell forward.

EGF-induced cell adhesion is brought about by enhanced formation of FAK⁺/Actin⁺ focal adhesion plaques, which is triggered by redistribution of activated FAK. EGF-induced formation of focal adhesions in serum-starved U373 cells was examined by multicolor immunofluorescence and confocal laser scanning microscopy using a murine monoclonal anti-FAK antibody (green fluorescence) and rhodamine-labeled phalloidin which stains actin (red fluorescence).

To evaluate the actin polymerization process, cells plated on poly-L-lysine-coated plates were first serum-starved to depolymerize the actin stress fibers. Subsequently, cells were stimulated with fetal bovine serum to induce *de novo* stress fiber formation.

Fluorescence microscopy

Immunofluorescence was used to study the effects of quinazoline derivatives on the formation of focal adhesion plaques and polymerization of actin. Cells (obtained and maintained as described for Example 6) were plated on poly-L-lysine-coated glass-bottom 35 mm Petri dishes (Mattek Corp., Ashland, MA) or fibronectin-coated cover slips and maintained in DMEM supplemented with 10% FBS for 24 hrs. The medium was removed and the cells were washed twice with serum-free DMEM and incubated in the same medium for 16 hours. Following this serum starvation, cells were incubated with varying concentrations of WHI-P131, WHI-P154 or vehicle (0.1% DMSO) for 4-16 hours at 37°C and then stimulated either with 250 ng/ml of human recombinant EGF or 10% FBS for 15, 30, 60, 120 or 180 minutes at 37°C. At the end of the EGF stimulation, cells were washed twice with PBS, fixed in 2% paraformaldehyde in PBS (pH 7.2), permeabilized and non-specific binding sites were blocked with 1.5% BSA and 0.1% triton X-100 in PBS for 30 minutes.

To detect the focal adhesion plaques and actin, cells were incubated with a mixture of a mouse monoclonal antibody directed against the focal adhesion kinase at a dilution of 1:100 and rhodamine-labeled phalloidin at a dilution of 1:1000 for 1 hour at room temperature. Cells were washed with PBS and incubated with a FITC-conjugated anti-mouse IgG (Amersham Corp., Arlington Heights, IL) for 1 hour (final dilution: 1:40). Cells were washed with PBS, counterstained with TOTO-3 (Molecular Probes, INC.) at a dilution of 1:1000 for 10 minutes at room temperature, washed again with PBS and the coverslips were counted with Vectashield (Vector Labs, Burlingame, CA). Subsequently, cells were viewed with a confocal laser scanning microscope (Bio-Rad MRC 1024) mounted in a Nikon Labhophot upright microscope. Digital images were saved on a Jaz disk and processed with Adobe Photoshop software (Adobe systems, Mountain View, CA). Prism software. Each experiment was repeated three times.

Results

As shown in **Figures 12A, 12A', 12B, 12B', 12C, 12C'**, a two-hour stimulation of serum-starved U373 cells with EGF resulted in a significant decrease of the diffuse perinuclear/cytoplasmic FAK staining accompanied by the emergence of focal adhesion plaques with high intensity FAK staining (bright green fluorescence). These FAK⁺ adhesion plaques showed a strong phalloidin staining (bright red fluorescence) confirming the colocalization of actin. Notably, preincubation of U373 cells with WHI-P154 (**Figure 12B**) or WHI-P131 (data not shown) at a 10 μ M concentration prevented the formation of FAK⁺/Actin⁺ focal adhesion plaques after EGF stimulation.

To evaluate the actin polymerization process, cells plated on poly-L-lysine-coated plates were first serum-starved to depolymerize the actin stress fibers. Subsequently, cells were stimulated with fetal bovine serum to induce *de novo* stress fiber formation. As shown in **Figure 13A-C**, a two-hour stimulation of serum-starved U373 cells with fetal bovine serum (10% v/v) resulted in a marked increase in polymerized actin stress fibers. Pretreatment of serum-starved U373 cells with WHI-P154 inhibited serum-induced actin polymerization (**Figure 13C**). Similar results were obtained with WHI-P131 but not with the unsubstituted dimethoxy quinazoline compound WHI-P258 (data not shown).

In summary, the data provided in Examples 6–9 demonstrate the effectiveness of substituted quinazolines in the inhibition of glioblastoma cell adhesion and migration, key factors for tumor cell metastasis. The most potent inhibitory agents were WHI–P154 and WHI–P131. Both compounds inhibited
5 adhesion and migration at micromolar concentration.

A complex network of intracellular molecules including receptor tyrosine kinases and Src family tyrosine kinases in cooperation with several extracellular factors such as substratum to which cells adhere and external factors, regulates the cell adhesion and motility (Finchman, V.J., and Frame, M.C., 1998,
10 *EMBO J.*, 17:81–92). Activation of integrin family adhesion receptors upon binding to specific extracellular matrix proteins has been shown to enhance the phosphorylation of integrins and activation of several intracellular signaling proteins including mitogen activated protein kinase, FAK, Src tyrosine kinases as well as p130^{cas}, talin, paxillin, and cortactin which were identified as substrates for the Src
15 tyrosine kinase (Cobb, B.S., Schaller, M.D., Leu, T.H., and Parsons, J.T., 1994, *Mol. Cell Biol.*, 14:147–155; Chen, Q., Lin, T.H., Der, C.J., Juliano, R.L., 1996, *J. Biol. Chem.*, 271:18122–18127; Klemke, R.L., Cai, S., Giannini, A.L., Gallagher, P.J., Lanerolle, P.D., and Cheresh, D.A., 1997, *J. Cell Biol.*, 137:481–492; Petch, L.A., Bockholt, S.M., Bouton, A., Parsons, J.T., Burridge, K., 1995, *J. Cell Sci.*,
20 108:1371–1379; Chrzanowska–Wodnicka, M., and Burridge, K., 1996, *J. Cell Biol.*, 133(6):1403–15; Miyamoto, S., Akiyama, S.K., Yamada, K.M., 1995, *Science*, 267:883–5; Miyamoto, S., Teramoto, H., Gutkind, J.S., and Yamada, K.M., 1996, *J. Cell Biol.*, 135:1633–1642; Chen, H.C., Appeddu, P.A., Parsons, J.T., Hildebrand, J.D., Schaller, M.D., and Guan, J.L., 1995, *J. Biol. Chem.*, 16995–16999).

25 Subsequently, the adhesion of cell is strengthened by redistribution of activated Src kinase and focal adhesion kinase to focal adhesions, recruitment and aggregation of activated intracellular proteins such as paxillin, talin, vinculin and clustering of integrins (Burridge, K., Fath, K., Kelly, G., and Turner, C., 1988, *Ann. Rev. Cell Biol.*, 4:487–525; Burridge, K., and Fath, K., 1989, *Bioessays*, 10:104–108;
30 Finchman, V.J., and Frame, M.C., 1998, *EMBO J.*, 17:81–92). FAK is also activated by the binding of certain growth factors to their receptors in a mechanism that is independent of integrin activation (Hatai, M., Hashi, H., Mogi, A., Soga, H., Yokota, J., Yaoi, Y., 1994, *FEBS Lett.*, 350:113–116; Ouwens, D.M., Mikkers, H.M., van der Zon, G.C., Stein Gerlach, M., Ullrich, A., Maassen, J.A., 1996,
35 *Biochem J.*, 318:609–614).

In experiments not shown here, WHI-P154 was found to be a potent inhibitor of the EGF-R tyrosine kinase as well as Src family tyrosine kinases (Liu and Uckun, manuscript in preparation). Therefore, it was initially postulated that the effects of WHI-P154 on glioblastoma cells was due to its tyrosine kinase inhibitory properties. Surprisingly, however, WHI-P79 and WHI-P131, which are equally potent inhibitors of the EGF-R and Src family tyrosine kinases were not as effective as WHI-P154 and the broad spectrum tyrosine kinase inhibitor genistein did not affect glioblastoma cell adhesion and motility at concentrations which abrogate the enzymatic activity of the EGF-R kinase and Src family tyrosine kinases. Thus, the effects of WHI-P154 on U373 cells cannot be explained by its tyrosine kinase inhibitory properties alone.

Example 10

WHI-P154 Inhibits Glioblastoma Cell Migration from Spheroids

U373 glioblastoma spheroids of 200 to 400 micrometers in diameter were treated with WHI-P154 in 0.1% DMSO at varied concentrations. Cells were incubated with the inhibitor or with control DMSO in the absence of inhibitor compound for two hours, and then transferred to fibronectin-coated coverslips. The spheroids were then incubated in DMEM containing WHI-P154 at 37° C for 48 hours.

As shown in Figures 14 A-14 D, treatment of glioblastoma spheroids with WHI-P154 significantly inhibited cell migration from the spheroid as compared with the untreated control and in a dose-dependent manner. The Figure shows the following dosage treatments: A: control; B: 2.5 μ M; C: 4.5 μ M; and D: 10 μ M.

Example 11

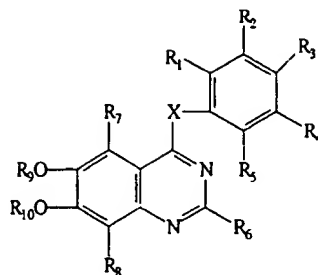
Cytotoxic Activity of WHI-P292

The novel compound WHI-P292 was assayed for cytotoxic activity in the MTT cell survival assay, as described above for Example 3. As shown in Figure 15, this compound demonstrated potent cytotoxic activity against glioblastoma cells, with an IC₅₀ of 38.22 μ M.

All publications, patents, and patent documents described herein are incorporated by reference as if fully set forth. The invention described herein may be modified to include alternative embodiments. All such obvious alternatives are within the spirit and scope of the invention, as claimed below.

We claim:

1. A compound comprising the formula:



wherein X is selected from the group consisting of HN, $R_{11}N$, S, O, CH_2 , and $R_{11}CH$;

R_{11} is H, alkyl having up to four carbon atoms, or acyl;

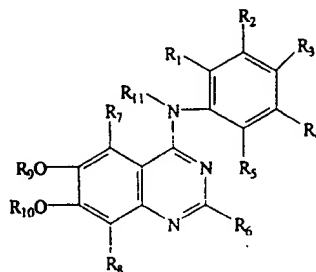
R_1 , R_2 , R_3 , R_4 and R_5 are the same or different and are each independently selected from the group consisting of H, OH, SH, NH_2 , NO_2 , halogen, alkoxy, acyl, and alkylthio, having up to four carbon atoms, and where one or more of R_1 - R_5 is OH, SH, or NH_2 ;

R_6 , R_7 and R_8 are the same or different and are each independently selected from the group consisting of H, OH, SH, NH_2 , NO_2 , acyl, alkoxy, alkylthio and halogens, having up to four carbon atoms; and

R_9 and R_{10} are the same or different and are each independently selected from the group consisting of H, alkyl or acyl, having up to four carbon atoms.

2. The compound of claim 1 wherein at least one of R_1 , R_2 , R_3 , R_4 or R_5 is OH.
3. The compound of claim 2 wherein at least one of R_1 , R_2 , R_3 , R_4 or R_5 is a halogen.
4. The compound of claim 1 wherein R_2 is OH.
5. The compound of claim 1 wherein R_2 is Cl, Br or F.

6. The compound of claim 1 wherein R_3 is OH.
7. The compound of claim 1 wherein R_4 is Br, Cl or F.
8. The compound of claim 1 wherein R_9 and R_{10} each comprise CH_3 .
9. The compound of claim 1 wherein R_{11} comprises H.
10. The compound of claim 1 wherein R_3 is OH and R_2 is halogen.
11. The compound of claim 1 wherein R_3 is OH and R_2 is Br.
12. A compound comprising the formula:



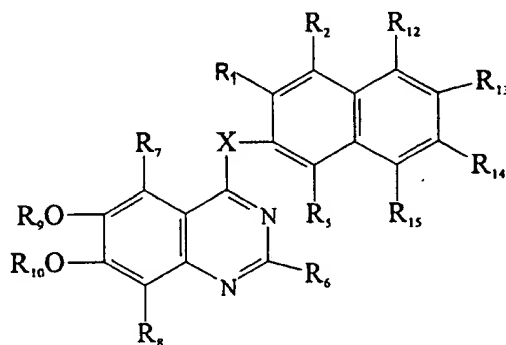
wherein R_{11} is H or alkyl having up to four carbon atoms;

R_1 , R_2 , R_3 , R_4 and R_5 are the same or different and are each independently selected from the group consisting of H, OH, SH, NH_2 , NO_2 , halogen, alkoxy, acyl, and alkylthio, having up to four carbon atoms, and where one or more of R_1 - R_5 is OH, SH, or NH_2 ;

R_6 , R_7 and R_8 are the same or different and are each independently selected from the group consisting of H, OH, SH, NH_2 , NO_2 , halogen, acyl, alkoxy, and alkylthio, having up to four carbon atoms; and

R_9 and R_{10} are the same or different and are each independently selected from the group consisting of H, alkyl or acyl, having up to four carbon atoms.

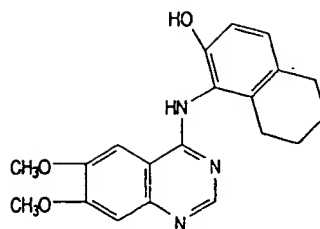
13. The compound of claim 12 wherein at least one of R_1 , R_2 , R_3 , R_4 or R_5 is OH, SH, or NH_2 .
14. The compound of claim 13 wherein at least one of R_1 , R_2 , R_3 , R_4 or R_5 is a halogen.
15. The compound of claim 12 wherein R_9 and R_{10} each comprise CH_3 .
16. The compound of claim 12 wherein R_3 is OH and R_2 is halogen.
17. The compound of claim 1, comprising a second ring fused to the phenyl ring via R_1 - R_5 .
18. The compound of claim 17 having the following structural formula:



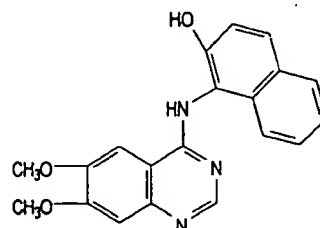
where one or more of R_1 , R_2 , R_5 , R_{12} , R_{13} , R_{14} , or R_{15} is OH, SH, or NH_2 .

19. The compound of claim 17, wherein one or more of R_1 , R_2 , R_5 , R_{12} , R_{13} , R_{14} , or R_{15} is OH.
20. The compound of claim 17 having the following structural formula:

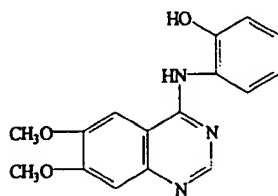
50



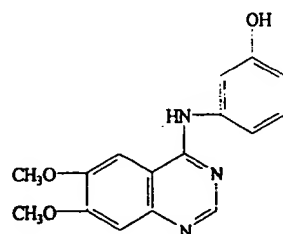
21. The compound of claim 17 having the following structural formula:



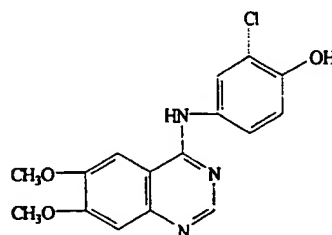
22. A compound having the structural formula:



23. A compound having the structural formula:

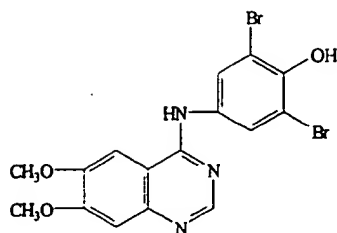


24. A compound having the structural formula:

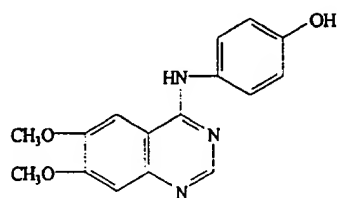


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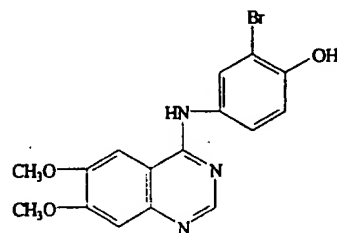
25. A compound having the structural formula:



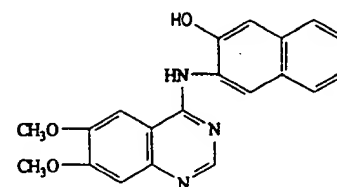
26. A compound having the structural formula:



27. A compound having the structural formula:



28. A compound having the structural formula:



29. A conjugate compound comprising a targeting moiety covalently bonded to a compound of claims 1.

30. The conjugate compound of claim 29 wherein the targeting moiety is an antibody, cytokine or growth factor.
31. The conjugate compound of claim 29 wherein the targeting moiety is an anti-cell surface receptor antibody.
32. The conjugate compound of claim 29 wherein the targeting moiety is EGF.
33. A conjugate compound comprising 4-(3'-Bromo-4'-hydroxy phenyl)-amino-6,7-dimethoxy quinazoline [WHI-P154) covalently bonded to EGF.
34. A conjugate compound comprising 4-(3'-Bromo-4'-hydroxy phenyl)-amino-6,7-dimethoxy quinazoline [WHI-P131) covalently bonded to EGF.
35. A conjugate compound comprising 4-(3'-Bromo-4'-hydroxy phenyl)-amino-6,7-dimethoxy quinazoline [WHI-P292) covalently bonded to EGF.
36. A composition comprising the compound of claim 1 and an isotonic carrier.
37. A composition comprising the compound of claim 12 and an isotonic carrier.
38. A composition comprising the conjugate compound of claim 29 and an isotonic carrier.
39. A method for inducing apoptosis of brain tumor cells, said method comprising contacting brain tumor cells with a compound of claim 1.
40. A method for inducing apoptosis of brain tumor cells, said method comprising contacting brain tumor cells with a compound of claim 12.

41. A method for inducing apoptosis of brain tumor cells, said method comprising contacting brain tumor cells with the conjugate of claim 29.

42. The method of claim 39 wherein said contacting is with an amount of 100 micromolar or greater of the compound.

43. A method for preventing adhesion or migration of brain tumor cells, said method comprising contacting brain tumor cells with the compound of claim 1.

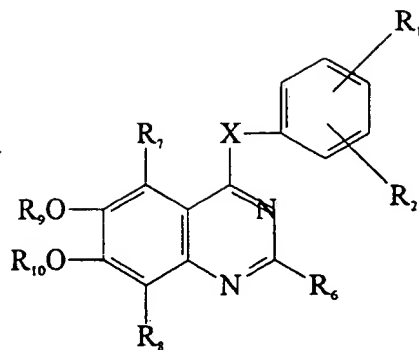
44. A method for preventing adhesion or migration of brain tumor cells, said method comprising contacting brain tumor cells with the compound of claim 12.

45. A method for preventing adhesion or migration of brain tumor cells, said method comprising contacting brain tumor cells with the conjugate of claim 29.

46. The method of claim 43 wherein said contacting is with an amount of 100 micromolar or less of the compound.

47. A method of treating brain tumor comprising administering to a brain tumor patient a conjugate compound comprising 4-(3'-bromo-4'-hydroxy phenyl)-amino-6, 7-dimethoxyquinazoline covalently bound to EGF.

48. A compound having the structural formula:



wherein X is selected from the group consisting of HN, $R_{11}N$, CH_2 , and $R_{11}CH$;

R_{11} is H, alkyl having up to four carbon atoms, or acyl wherein acyl is $-C(OR)$ and R is H or alkyl having up to four carbon atoms;

R_1 is halogen;

R_2 is OH;

R_6 , R_7 , and R_8 are the same or different and are each independently selected from the group consisting of H, OH, SH, NH_2 , NO_2 , halogen, alkoxy having up to four carbon atoms, alkylthio having up to four carbon atoms, and acyl wherein acyl is $-C(OR)$ and R is H or alkyl having up to four carbon atoms; and

R_9 and R_{10} are the same or different and are each independently selected from the group consisting of H, alkyl having up to four carbon atoms, and acyl wherein acyl is $-C(OR)$ and R is H or alkyl having up to four carbon atoms.

49. The compound according to claim 48, wherein X is HN.

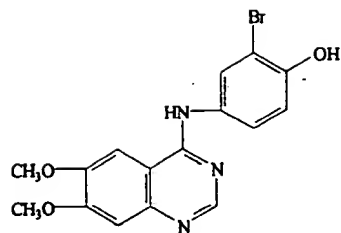
50. The compound according to claim 48, wherein R_1 is Br.

51. The compound according to claim 48, wherein R_1 is Cl.

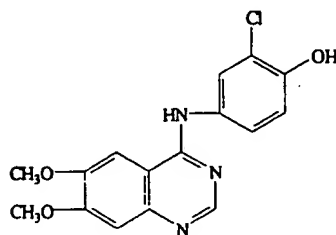
52. The compound according to claim 48, wherein R_1 is F.

53. A compound having the structural formula:

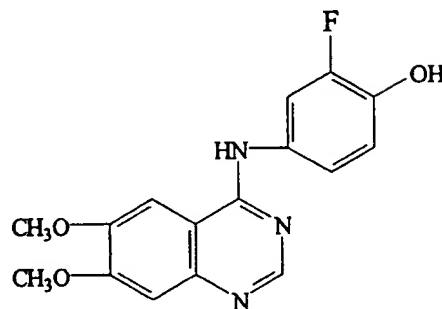
55



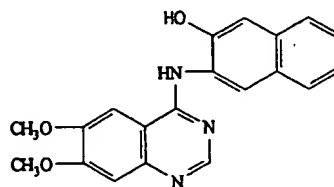
54. A compound having the structural formula:



55. A compound having the structural formula:



56. A compound having the structural formula:



57. A composition comprising the compound according to claim 48 and an isotonic carrier.

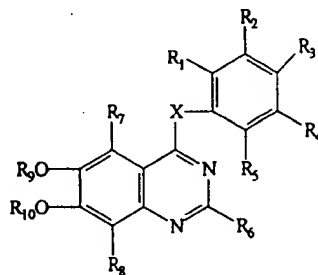
58. A composition comprising the compound according to claim 53 and an isotonic carrier.

59. A composition comprising the compound according to claim 54 and an isotonic carrier.

60. A composition comprising the compound according to claim 55 and an isotonic carrier.

61. A composition comprising the compound according to claim 56 and an isotonic carrier.

62. A method for inducing apoptosis of brain tumor cells, said method comprising contacting brain tumor cells with an effective apoptosis-inducing amount of a compound comprising the formula:



wherein R_1 , R_2 , R_3 , R_4 and R_5 are the same or different and are each independently H, OH, SH, NH_2 , NO_2 , halogen, alkoxy having up to four carbon atoms, alkylthio having up to four carbon atoms, or acyl wherein acyl is $-C(OR)$ and R is H or alkyl having up to four carbon atoms, or any two of R_1 - R_5 may be taken together to form a saturated, unsaturated, or aromatic ring, and where one or more of R_1 - R_5 is OH;

X is selected from the group consisting of HN, $R_{11}N$, S, O, CH_2 , and $R_{11}CH$;

R_{11} is H, alkyl having up to four carbon atoms, or acyl wherein acyl is $-C(OR)$ and R is H or alkyl having up to four carbon atoms;

R_6 , R_7 , and R_8 are the same or different and are each independently selected from the group consisting of H, OH, SH, NH_2 , NO_2 , halogen, alkoxy having up to four carbon atoms, alkylthio having up to four carbon atoms, and acyl wherein acyl is $-C(OR)$ and R is H or alkyl having up to four carbon atoms; and

R_9 and R_{10} are the same or different and are each independently selected from the group consisting of H, alkyl having up to four carbon atoms, and acyl wherein acyl is $-C(OR)$ and R is H or alkyl having up to four carbon atoms.

63. The method according to claim 62, wherein one or more of R_1 - R_5 is halogen.

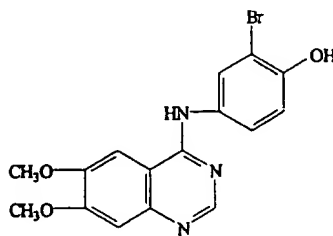
64. The method according to claim 63, wherein one or more of R_1 - R_5 is Br.

65. The method according to claim 63, wherein one or more of R_1 - R_5 is Cl.

66. The method according to claim 63, wherein one or more of R_1 - R_5 is F.

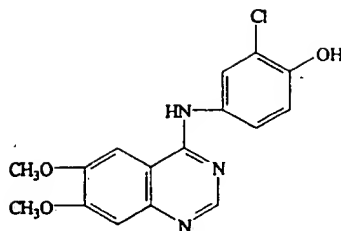
67. The method according to claim 62, wherein X is HN.

68. The method according to claim 62, wherein the compound has the formula:

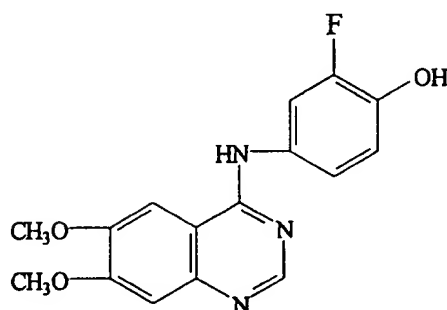


69. The method according to claim 62, wherein the compound has the formula:

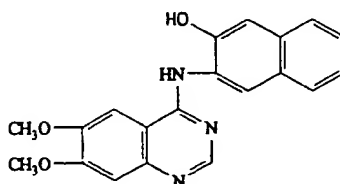
58



70. The method according to claim 62, wherein the compound has the formula:

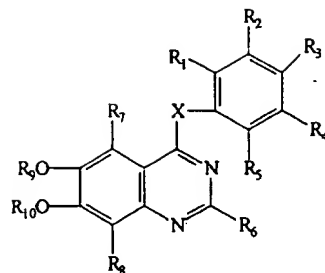


71. The method according to claim 62, wherein the compound has the formula:



72. The method of claim 62, wherein said contacting is with an amount of 100 micromolar or greater of the compound.

73. A method for preventing adhesion or migration of brain tumor cells, said method comprising contacting brain tumor cells with a compound comprising the formula:



wherein R_1 , R_2 , R_3 , R_4 and R_5 are the same or different and are each independently H, OH, SH, NH_2 , NO_2 , halogen, alkoxy having up to four carbon atoms, alkylthio having up to four carbon atoms, or acyl wherein acyl is $-\text{C}(\text{OR})$ and R is H or alkyl having up to four carbon atoms, or any two of R_1 - R_5 may be taken together to form a saturated, unsaturated, or aromatic ring, and where one or more of R_1 - R_5 is OH;

X is selected from the group consisting of HN, $R_{11}\text{N}$, S, O, CH_2 , and $R_{11}\text{CH}$;

R_{11} is H, alkyl having up to four carbon atoms, or acyl wherein acyl is $-\text{C}(\text{OR})$ and R is H or alkyl having up to four carbon atoms;

R_6 , R_7 , and R_8 are the same or different and are each independently selected from the group consisting of H, OH, SH, NH_2 , NO_2 , halogen, alkoxy having up to four carbon atoms, alkylthio having up to four carbon atoms, and acyl wherein acyl is $-\text{C}(\text{OR})$ and R is H or alkyl having up to four carbon atoms; and

R_9 and R_{10} are the same or different and are each independently selected from the group consisting of H, alkyl having up to four carbon atoms, and acyl wherein acyl is $-\text{C}(\text{OR})$ and R is H or alkyl having up to four carbon atoms.

74. The method according to claim 73, wherein one or more of R_1 - R_5 is halogen.

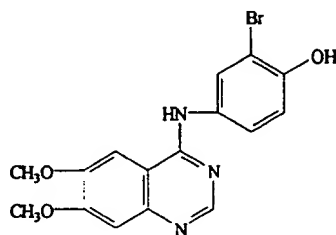
75. The method according to claim 73, wherein one or more of R_1 - R_5 is Br.

76. The method according to claim 73, wherein one or more of R_1 - R_5 is Cl.

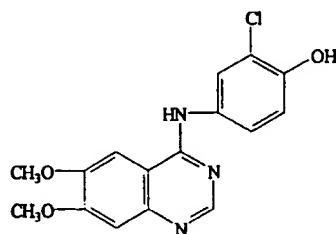
77. The method according to claim 73, wherein one or more of R_1 - R_5 is F.

78. The method according to claim 73, wherein X is HN.

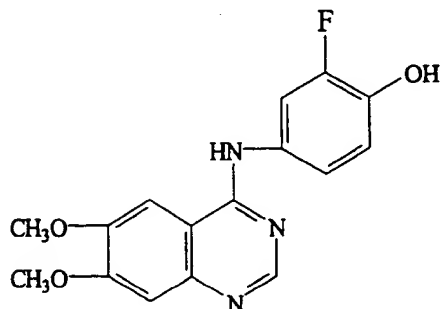
79. The method according to claim 73, wherein the compound has the formula:



80. The method according to claim 73, wherein the compound has the formula:

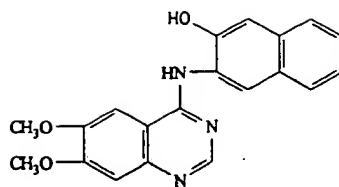


81. The method according to claim 73, wherein the compound has the formula:



82. The method according to claim 73, wherein the compound has the formula:

61



83. The method according to claim 73, wherein said contacting is with an amount of 100 micromolar or less of the compound.

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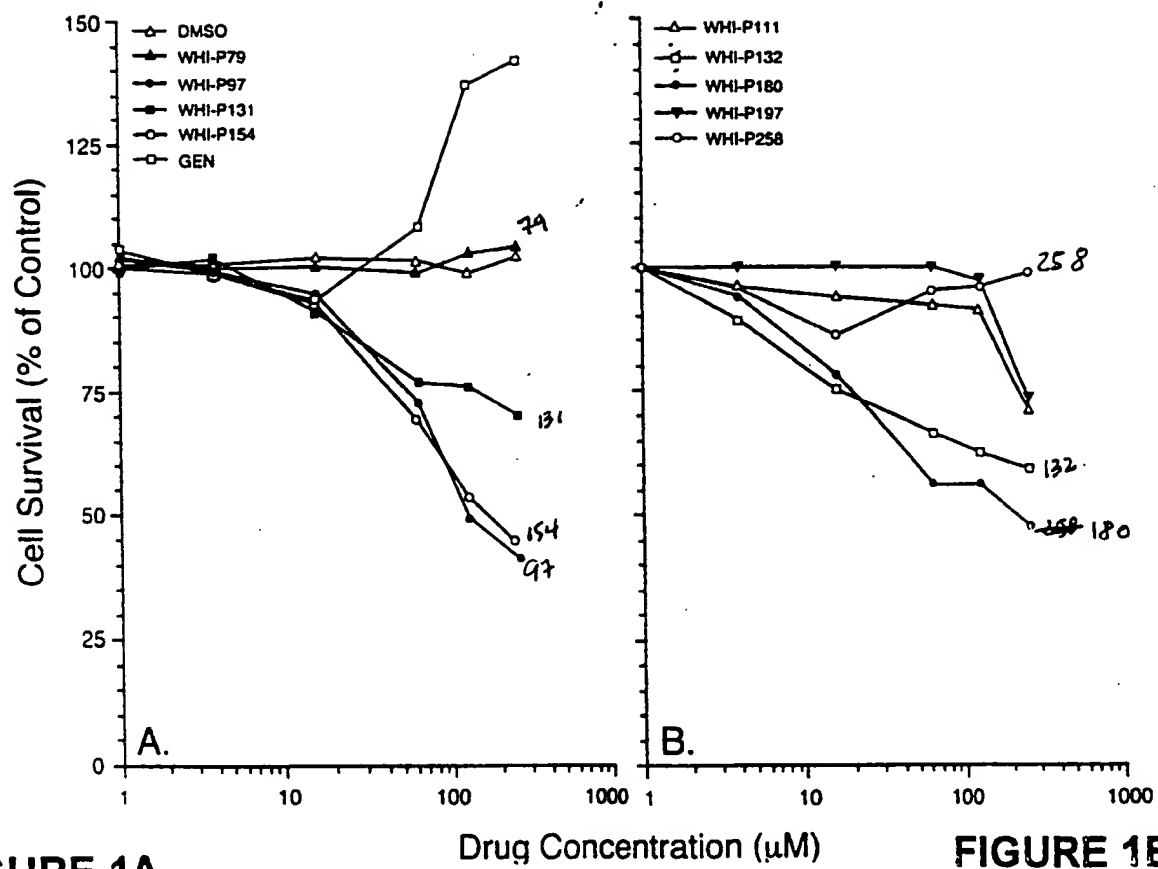


FIGURE 1A

FIGURE 1B

FIGURE 2A

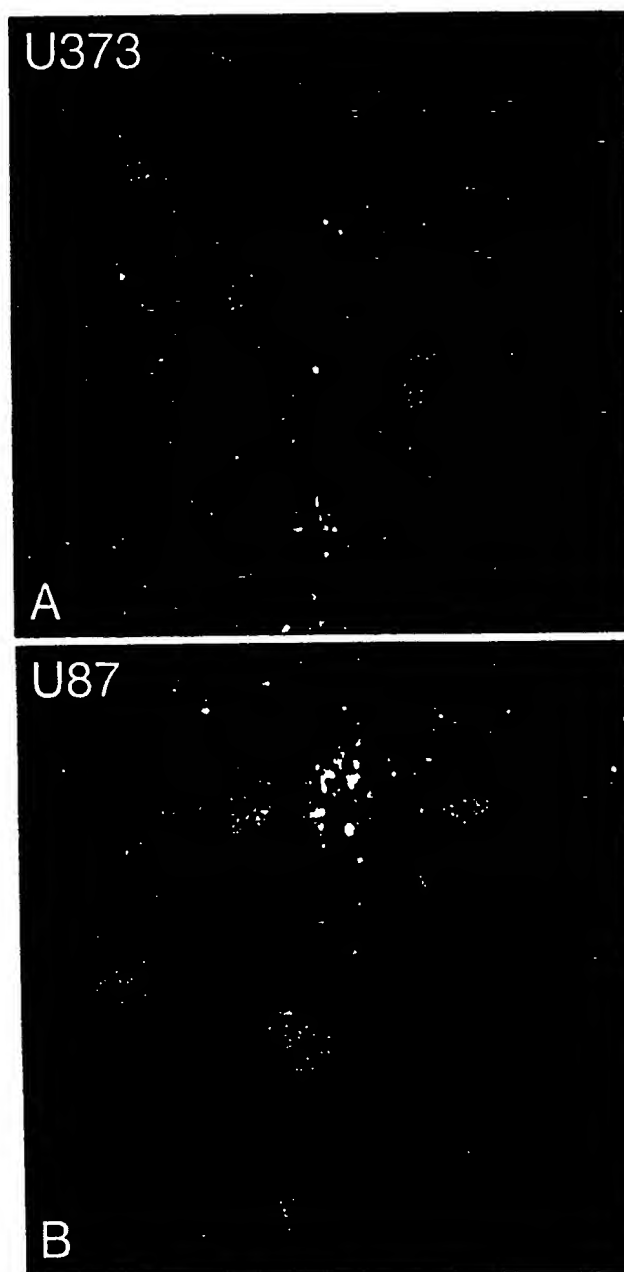
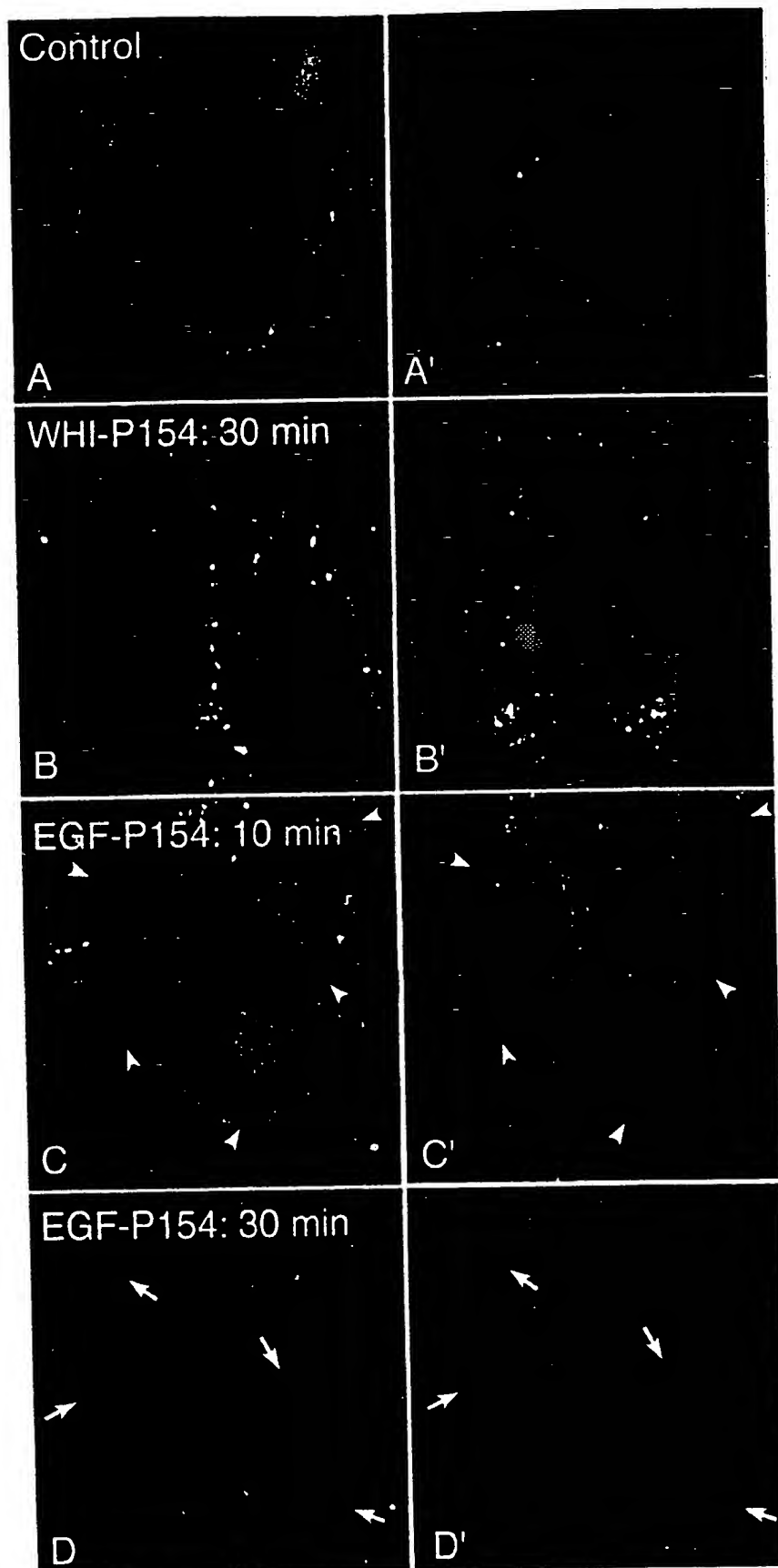
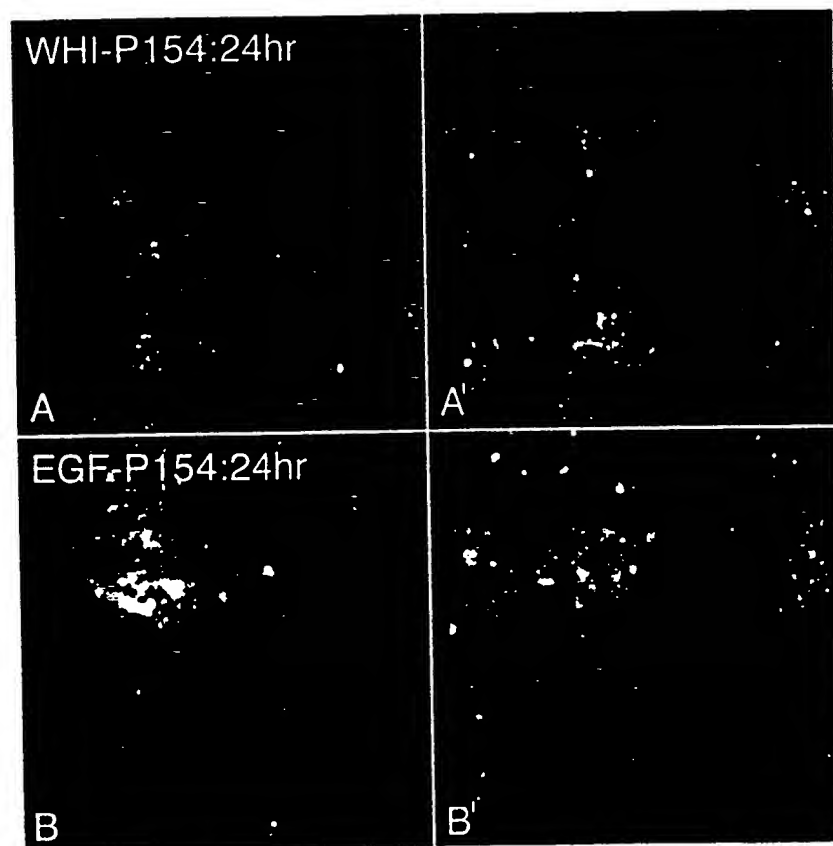


FIGURE 2B

FIGURE 3

**FIGURE 4.**

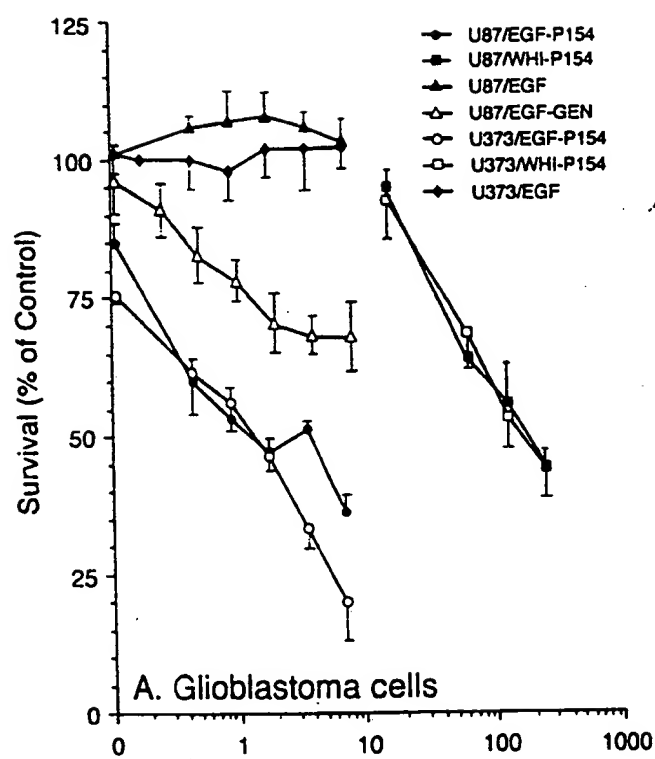


FIGURE 5A

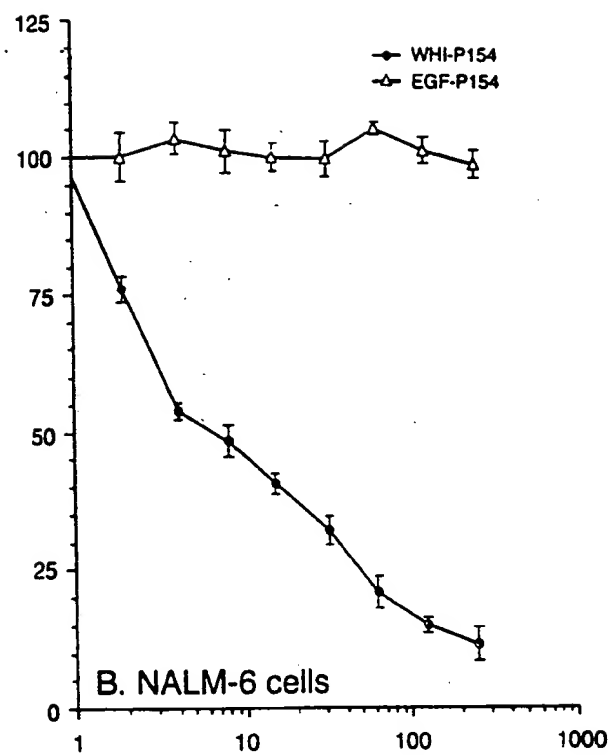
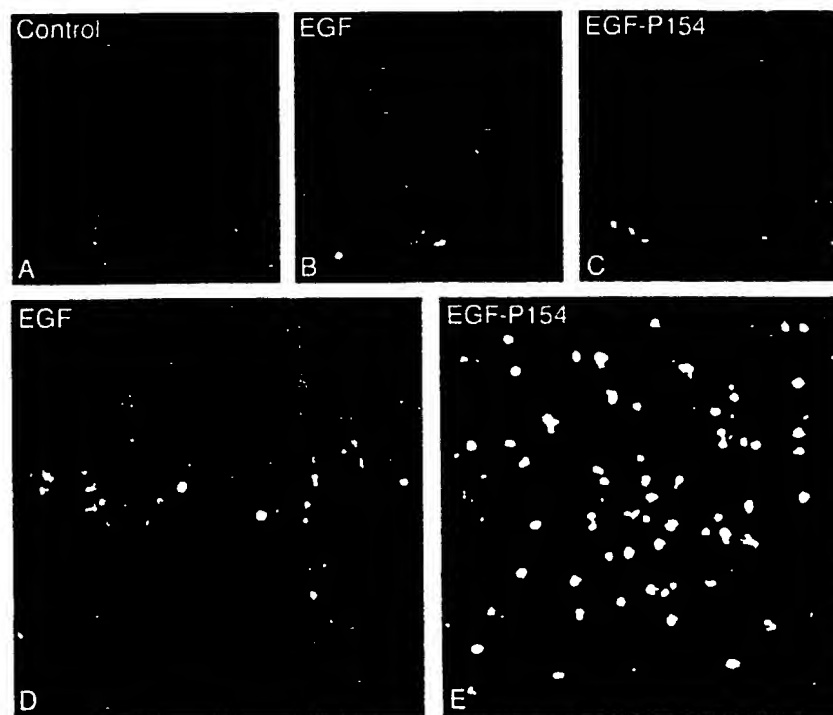


FIGURE 5B

FIGURE 6.

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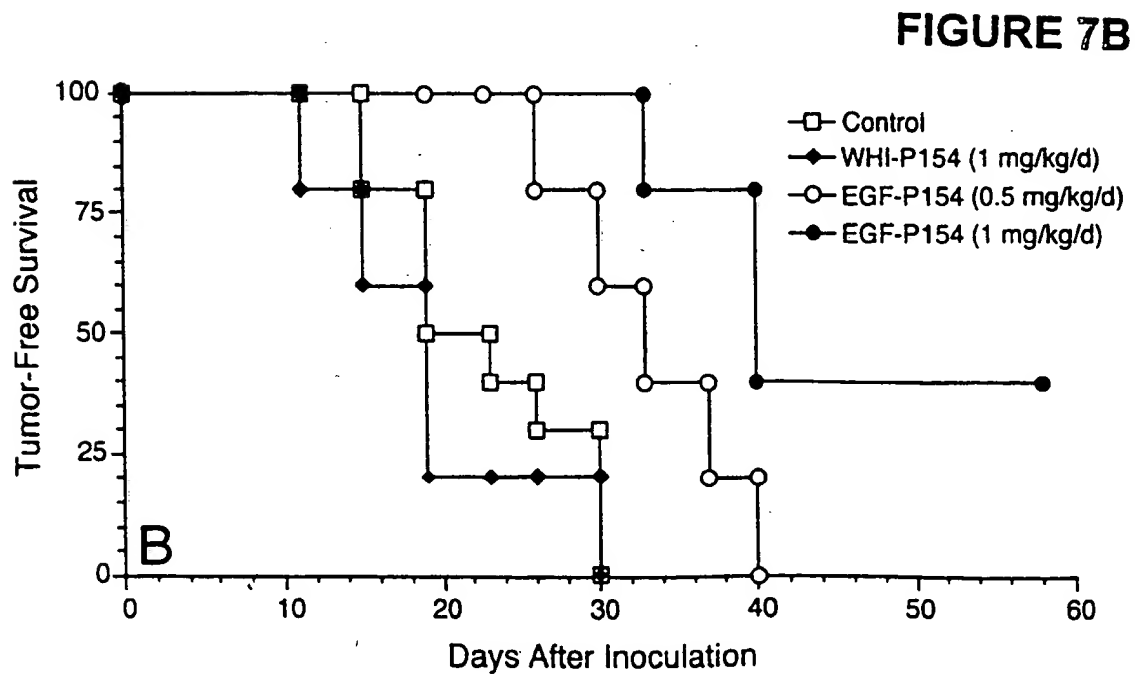
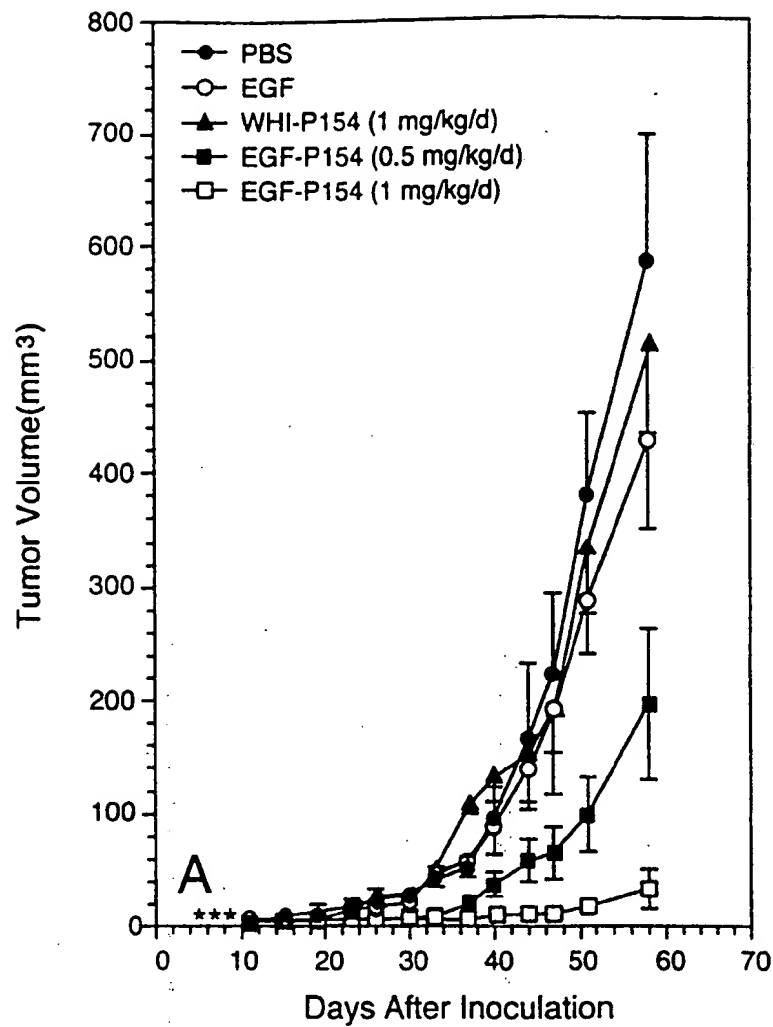
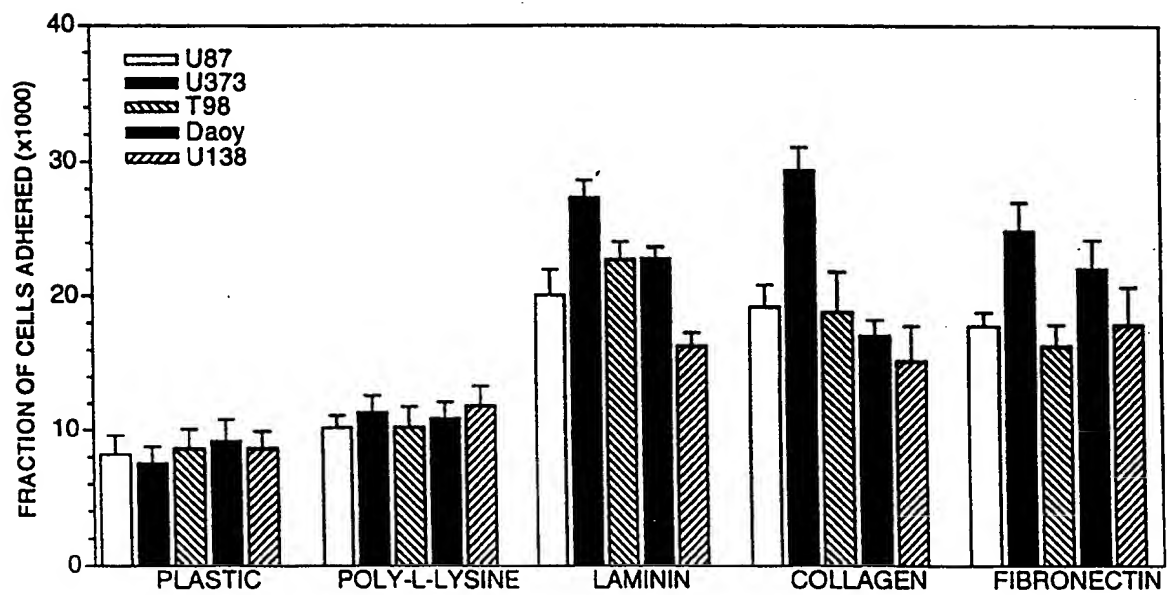


FIGURE 8



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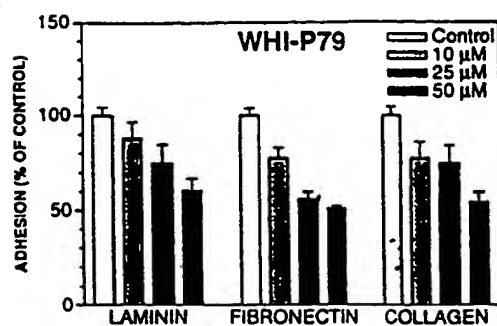


FIGURE 9A

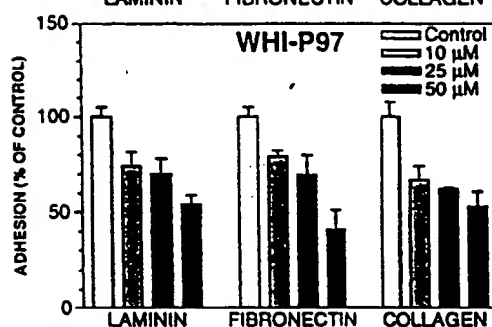


FIGURE 9B

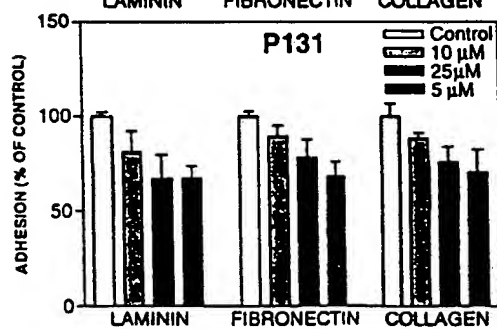


FIGURE 9C

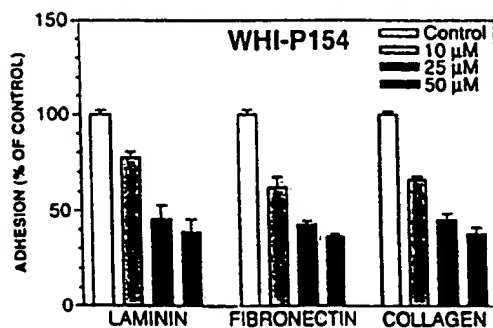
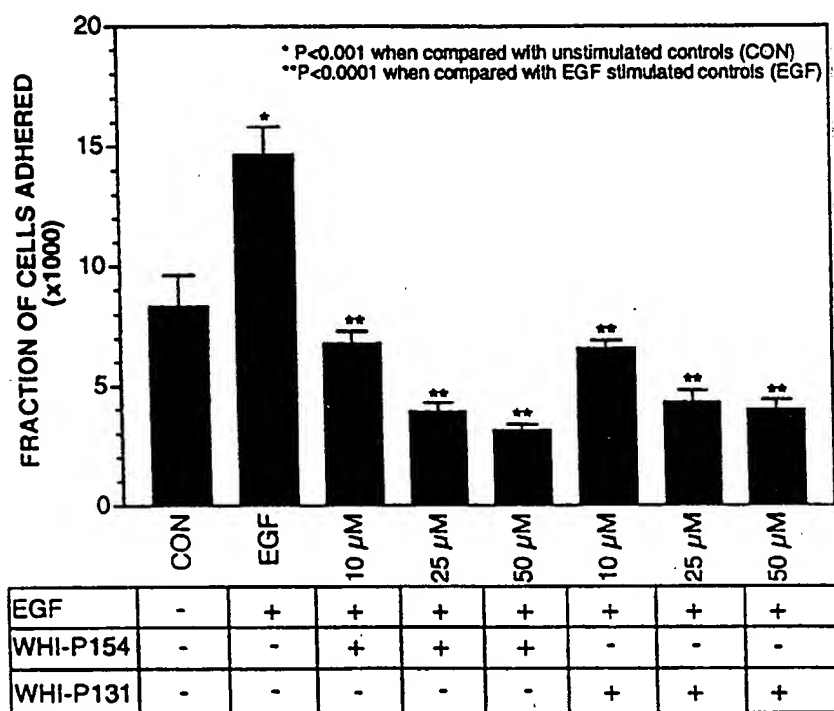


FIGURE 9D

FIGURE 10



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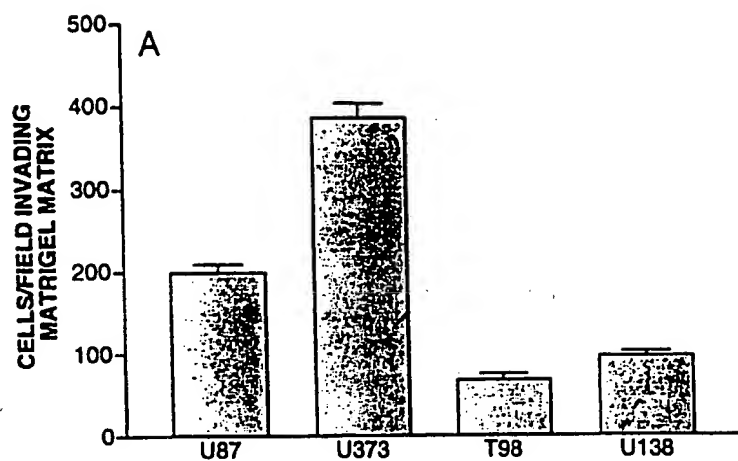
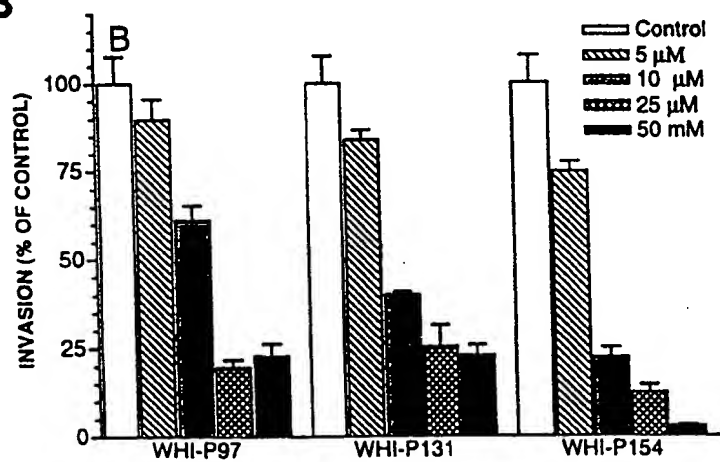
FIGURE 11A**FIGURE 11B**

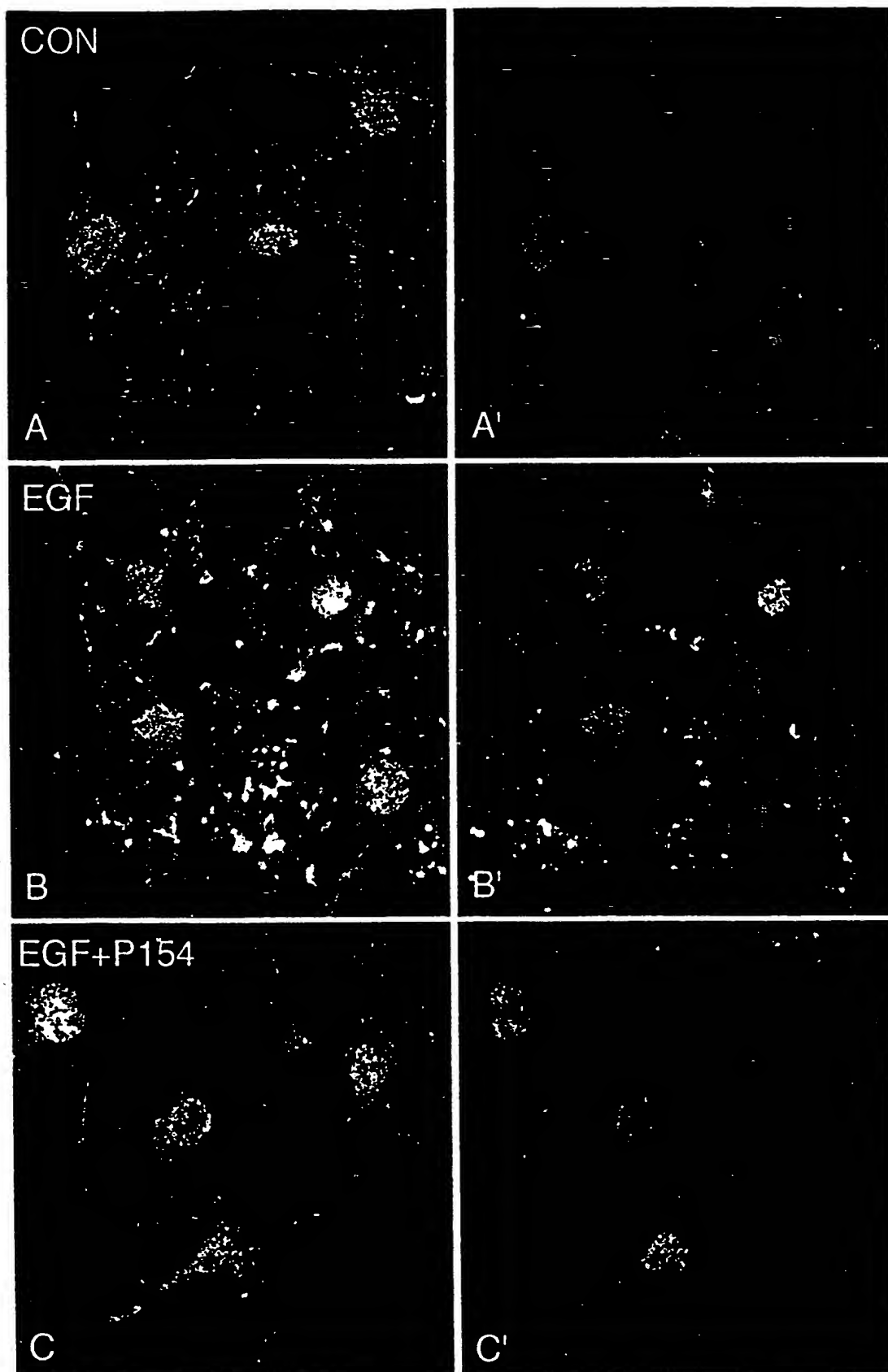
FIGURE 12

FIGURE 13

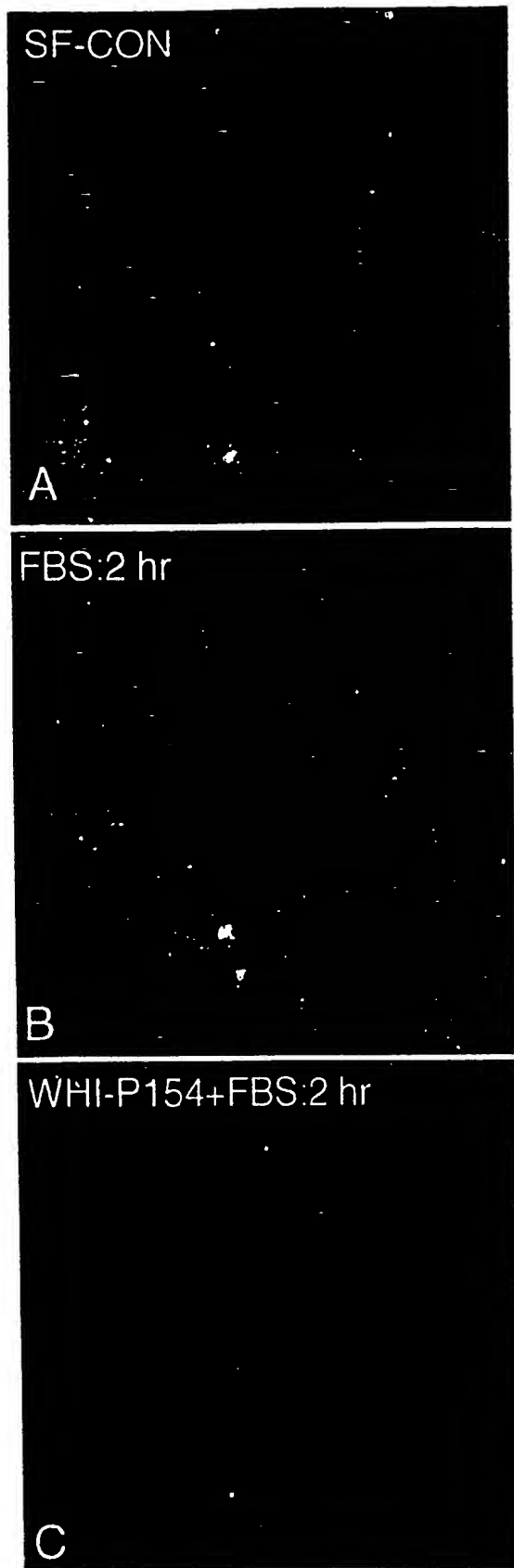


FIGURE 14

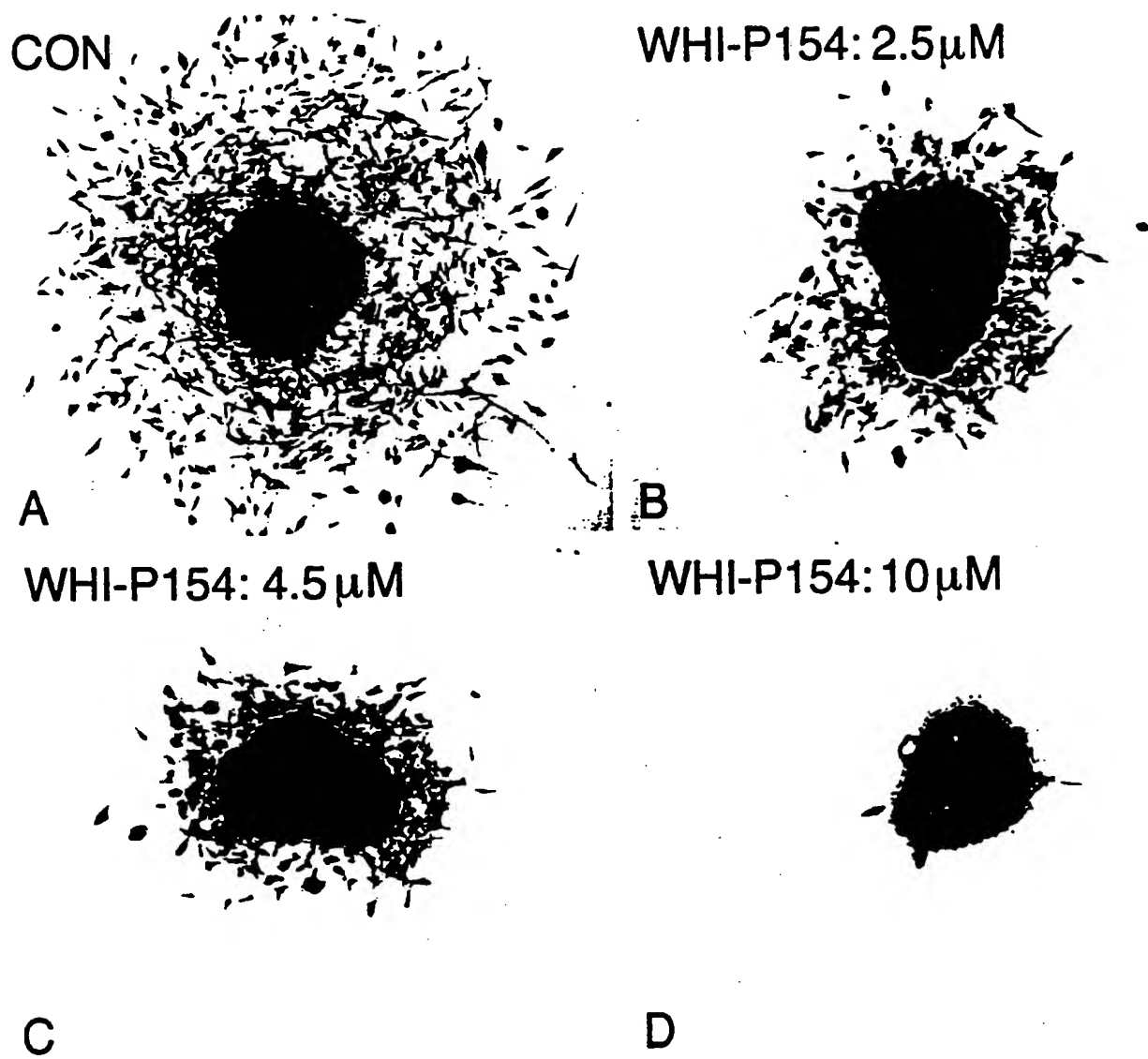
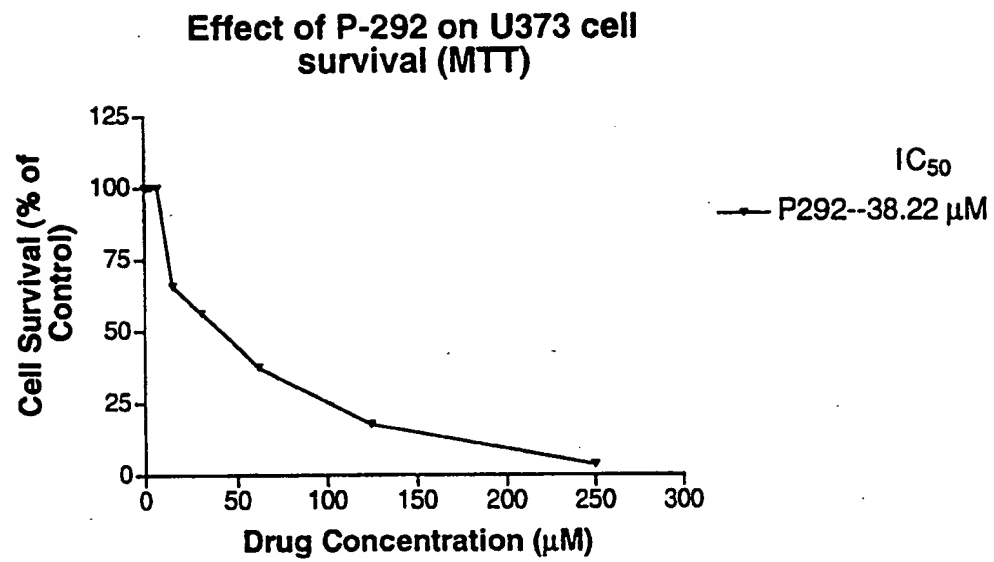


FIGURE 15

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/11767

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07D239/94 C07D239/88 C07D239/74 A61K31/505

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 566 226 A (ZENECA) 20 October 1993 (1993-10-20) claims; example 69 ---	1-13, 18, 26, 62, 73
X	WO 97 32856 A (ZENECA) 12 September 1997 (1997-09-12) claims; examples 2, 3 ---	1-3, 8, 12, 18, 62, 73
X	WO 97 30035 A (ZENECA) 21 August 1997 (1997-08-21) claims; examples 1-5, 9, 11, 12 ---	1-9, 12, 25, 62, 73
X	WO 96 15118 A (ZENECA) 23 May 1996 (1996-05-23) claims ---	1-3, 8, 12, 18, 62, 73
-/-		

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"Z" document member of the same patent family

Date of the actual completion of the international search

24 August 1999

Date of mailing of the international search report

06/09/1999

Name and mailing address of the ISA

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Fax: (+31-70) 340-3016

Authorized officer

Francois, J

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/11767

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 95 15758 A (RHONE-POULENC) 15 June 1995 (1995-06-15) claims	1-3,8, 12,18, 62,73
P,X	R. K.NARLA ET AL.: "4-(3'-BROMO-4'-HYDROXYPHENYL)AMINO-6,7-DIM ETHOXYQUINAZOLINE:" CLINICAL CANCER RESEARCH., vol. 4, no. 6, June 1998 (1998-06), pages 1405-1414, XP002113201 THE AMERICAN ASSOCIATION FOR CANCER RESEARCH., US ISSN: 1078-0432 page 1404 - page 1407	1-12, 23-27, 62,73

INTERNATIONAL SEARCH REPORT

national application No.

PCT/US 99/ 11767

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 47
because they relate to subject matter not required to be searched by this Authority, namely:
Remark: Although claims 47
is directed to a method of treatment of the human/animal
body, the search has been carried out and based on the alleged
effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such
an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all
searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment
of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report
covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is
restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No

PCT/US 99/11767

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 566226	A	20-10-1993	AT 130000 T	15-11-1995
			AU 661533 B	27-07-1995
			AU 3101093 A	22-07-1993
			CA 2086968 A,C	21-07-1993
			CZ 282038 B	16-04-1997
			DE 69300754 D	14-12-1995
			DE 69300754 T	28-03-1996
			DK 566226 T	18-03-1996
			ES 2078798 T	16-12-1995
			FI 930208 A	21-07-1993
			GR 3018143 T	29-02-1996
			HK 36497 A	04-04-1997
			HU 9500185 A	28-07-1995
			MX 9300277 A	30-06-1994
			NO 301541 B	10-11-1997
			NZ 245662 A	26-09-1995
			SK 1693 A	09-09-1993
			US 5457105 A	10-10-1995
			US 5616582 A	01-04-1997
			ZA 9300015 A	20-07-1993
			JP 6073025 A	15-03-1994
WO 9732856	A	12-09-1997	AU 1866497 A	22-09-1997
			CA 2244897 A	12-09-1997
			CN 1212684 A	31-03-1999
			EP 0885198 A	23-12-1998
			NO 984085 A	04-09-1998
			ZA 9701747 A	27-08-1998
WO 9730035	A	21-08-1997	AU 1729097 A	02-09-1997
			CA 2242425 A	21-08-1997
			CN 1211239 A	17-03-1999
			CZ 9802535 A	11-11-1998
			EP 0880508 A	02-12-1998
			NO 983687 A	13-08-1998
			PL 328310 A	18-01-1999
WO 9615118	A	23-05-1996	AT 175962 T	15-02-1999
			AU 703328 B	25-03-1999
			AU 3813095 A	06-06-1996
			CA 2200871 A	23-05-1996
			DE 69507505 D	04-03-1999
			DE 69507505 T	02-06-1999
			EP 0790986 A	27-08-1997
			ES 2128092 T	01-05-1999
			FI 971970 A	07-05-1997
			JP 10508616 T	25-08-1998
			NO 972152 A	12-05-1997
			NZ 294917 A	25-02-1999
			US 5821246 A	13-10-1998
			ZA 9509572 A	13-05-1996
WO 9515758	A	15-06-1995	US 5480883 A	02-01-1996
			US 5710158 A	20-01-1998
			AU 1305095 A	27-06-1995
			EP 0871448 A	21-10-1998
			SG 54172 A	16-11-1998
			US 5795889 A	18-08-1998

INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No

PCT/US 99/11767

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9515758 A		US 5646153 A	08-07-1997
		US 5721237 A	24-02-1998
		US 5714493 A	03-02-1998
		US RE36256 E	20-07-1999
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